# GenCore version 5.1.6 Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 13, 2003, 04:37:24; Search time 1703.81 Seconds

(without alignments)

7460.491 Million cell updates/sec

5

Title: US-09-852-261-5

Perfect score: 523

Sequence: 1 ggaccggagacgctctgcgg.....aaatacacaagtaaacattc 523

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

8:

Maximum Match 100%

Listing first 45 summaries

Database : EST:\*

1: em\_estba:\*
2: em\_esthum:\*
3: em\_estin:\*
4: em\_estmu:\*
5: em\_estov:\*
6: em\_estpl:\*
7: em\_estro:\*

9: gb\_est1:\*
10: gb\_est2:\*

em htc:\*

11: gb\_htc:\*

12: gb\_est3:\*

13: gb\_est4:\* 14: gb est5:\*

15: em\_estfun:\*

16: em\_estom:\*
17: em\_gss\_hum:\*

18: em\_gss\_inv:\*

19: em\_gss\_pln:\*

20: em\_gss\_vrt:\*

21: em\_gss\_fun:\*
22: em gss mam:\*

23: em\_gss\_mus:\*

24: em\_gss\_pro:\*

25: em\_gss\_rod:\*

26: em\_gss\_phg:\*

27: em gss vrl:\*

28: gb\_gss1:\*
29: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

						SUMMAN	מם
			ક				
Res	sult		Query				D
	No.	Score	Match	Length	DB	ID	Description
c	1	364.8	69.8	558	9	AI503976	AI503976 vm43d08.x
С	2	363	69.4	623	9	AW146128	AW146128 um37e10.x
- C	3	348.2	66.6	549	9	AI169253	AI169253 EST215088
C	4	347	66.3	558	9	AI265629	AI265629 uj04b07.x
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	6	339.2	64.9	816	9	AI119218	AI119218 ue94h02.y
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	8	334.4	63.9	796	14	CB959991	CB959991 AGENCOURT
c	9	322.2	61.6	499	9	AW495481	AW495481 UI-M-BH3-
c	10	320.8	61.3	642	9	AI876493	AI876493 uj59b10.x
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c	13	309.2	59.1	468	9	AI169770	AI169770 EST215669
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	29 29	263.8	50.4	799	9	AI314558	AI314558 uj48d07.y
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DEFINITION
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ACCESSION
            AI503976
            AT503976.1 GI:4401827
VERSION
            EST.
KEYWORDS
            Mus musculus (house mouse)
SOURCE
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              (bases 1 to 558)
REFERENCE
            Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T.,
  AUTHORS
            Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person
            ,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter
            ,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
            Waterston, R. and Wilson, R.
            The WashU-NCI Mouse EST Project 1999
  TITLE
            Unpublished
  JOURNAL
            Contact: Marra M/WashU-NCI Mouse EST Project 1999
COMMENT
            Washington University School of Medicine
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: mouseest@watson.wustl.edu
            This clone is available royalty-free through LLNL; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            MGI:565223
            This clone was previously sequenced on the 5' end only, this new
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                     Location/Qualifiers
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ORIGIN

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                                     82: Indels
                                                 7: Gaps
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       Qy
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         preproinsulin-like growth factor IB (MOUSE);, mRNA sequence.
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VERSION
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KEYWORDS
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SOURCE
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REFERENCE
         Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T.,
  AUTHORS
         Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person
```

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,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter
           ,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
           Waterston, R. and Wilson, R.
           The WashU-NCI Mouse EST Project 1999
 TITLE
           Unpublished
 JOURNAL
           Contact: Marra M/WashU-NCI Mouse EST Project 1999
COMMENT
           Washington University School of Medicine
           4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
           Tel: 314 286 1800
           Fax: 314 286 1810
           Email: mouseest@watson.wustl.edu
           This clone is available royalty-free through LLNL; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
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                   double-stranded cDNA was ligated to a DraIII adaptor
                   [TGTTGGCCTACTGG], digested and cloned into distinct DraIII
                   sites of the pME18S-FL3 vector (5' site CACTGTGTG, 3' site
                   CACCATGTG). XhoI should be used to isolate the cDNA
                   insert. Size selection was performed to exclude fragments
                   <1.5kb. Library constructed by Dr. Sumio Sugano
                    (University of Tokyo Institute of Medical Science).
                   Custom primers for sequencing: 5' end primer
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                                        191 t
                                                  1 others
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                       138 c
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  Query Match
                        81.9%; Pred. No. 2.5e-80;
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REFERENCE
                      Lee, N.H., Glodek, A., Chandra, I., Mason, T.M., Quackenbush, J.,
    AUTHORS
                      Kerlavage, A.R. and Adams, M.D.
                      Rat Genome Project: Generation of a Rat EST (REST) Catalog & Rat
    TITLE
                      Gene Index
    JOURNAL
                      Unpublished
                      On Oct 6, 1998 this sequence version replaced gi:3705561.
COMMENT
                      Other ESTs: TC50779
                      Contact: Lee, NH
                      The Institute for Genomic Research
                      9712, Medical Center Drive, Rockville, MD 20850, USA
                      Tel: (301)-838-3529
                      Fax: (301)-838-0208
                      Email: nhlee@tigr.org
                      Seq primer: M13-21.
                                      Location/Qualifiers
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DEFINITION
         IMAGE:1890901 3' similar to gb:X04482 Mouse mRNA for
         preproinsulin-like growth factor IB (MOUSE);, mRNA sequence.
ACCESSION
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KEYWORDS
            EST.
            Mus musculus (house mouse)
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            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
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               (bases 1 to 558)
REFERENCE
            Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
 AUTHORS
            Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
            Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,
            Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and
            Waterston, R.
            The WashU-HHMI Mouse EST Project
 TTTLE
            Unpublished
  JOURNAL
            Contact: Marra M/Mouse EST Project
COMMENT
            WashU-HHMI Mouse EST Project
            Washington University School of MedicineP
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: mouseest@watson.wustl.edu
            This clone is available royalty-free through LLNL; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            MGI:975225
            Seg primer: custom primer used
            High quality sequence stop: 495.
                     Location/Oualifiers
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                     /dev stage="adult"
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                     /note="Organ: liver; Vector: pME18S-FL3; Site_1: DraIII
                     (CACTGTGTG); Site_2: DraIII (CACCATGTG); 1st strand cDNA
                     was primed with an oligo(dT) primer
                     [ATGTGGCCTTTTTTTTTTTTTTTT]; double-stranded cDNA was
                     ligated to a DraIII adaptor [TGTTGGCCTACTGG], digested
                     and cloned into distinct DraIII sites of the pME18S-FL3
                     vector (5' site CACTGTGTG, 3' site CACCATGTG). XhoI should
                     be used to isolate the cDNA insert. Size selection was
                     performed to exclude fragments <1.5kb. Library
                     constructed by Dr. Sumio Sugano (University of Tokyo
                     Institute of Medical Science). Custom primers for
                     sequencing: 5' end primer CTTCTGCTCTAAAAGCTGCG and 3' end
                     primer CGACCTGCAGCTCGAGCACA."
BASE COUNT
                106 a
                         135 c
                                  156 g
                                            161 t
ORIGIN
                          66.3%; Score 347; DB 9; Length 558;
  Query Match
                          82.0%; Pred. No. 2.5e-76;
  Best Local Similarity
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0; Mismatches

Matches 414; Conservative

85; Indels

6; Gaps

1;

AI265629.1 GI:3883787

VERSION

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1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGAGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
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Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
           446 AGGGGCTTTTACTTCAACAAGCCCACAGGCTATGGCTCCAGCATTCGGAGGGCACCTCAG 387
Db
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Qу
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Db
        Qу
           326 TGTGCCCCACTGAAGCCTACAAAAGCAGCCCGCTCTATCCGTGCCCAGCGCCACACTGAC 267
Db
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Qy
           Db
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Qу
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Qу
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Db
        421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA-----AAAAATAAGTTTGATC 474
Qу
                         11111 1 11 111 111
                                              +1 +1 +1 +1 +1
                     1
           | \cdot |
        86 CTGCTTGAGCAACCTGCAAAACATCGAAACACCTACCAAATAACAATAATAAGTCCAATA 27
Db
        475 ACATTTCAAAGATGGCATTTCCCCC 499
Qy
           11111111111
                         -1111111
        26 ACATTACAAAGATGGGCATTTCCCC 2
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         CD373004
         UI-R-GRO-csv-j-17-0-UI.rl UI-R-GRO Rattus norvegicus cDNA clone
DEFINITION
         UI-R-GRO-csv-j-17-0-UI 5', mRNA sequence.
         CD373004
ACCESSION
         CD373004.1 GI:31157094
VERSION
         EST.
KEYWORDS
SOURCE
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 ORGANISM
         Rattus norvegicus
         Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
         Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
         Rattus.
REFERENCE
            (bases 1 to 614)
         Bonaldo, M.F., Lennon, G. and Soares, M.B.
 AUTHORS
         Normalization and subtraction: two approaches to facilitate gene
 TITLE
         discovery
         Genome Res. 6 (9), 791-806 (1996)
  JOURNAL
 MEDLINE
         97044477
  PUBMED
         8889548
```

```
Contact: Soares, MB
COMMENT
           Coordinated Laboratory for Computational Genomics
           University of Iowa
           375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA
           Tel: 319 335 8250
           Fax: 319 335 9565
           Email: bento-soares@uiowa.edu
           Tissue Procurement: James Lin, University of Iowa
            cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
            cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
            DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
            Clone Distribution: Distribution information can be found at
           http://genome.uiowa.edu/distribution/rat.html
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                   /lab host="DH10B (Life Technologies) (T1 phage resistant)"
                   /clone_lib="UI-R-GR0"
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                   tissue(s): rat whole embryo 13dpc. The library was
                   constructed according to Bonaldo, Lennon and Soares,
                   Genome Research, 6:791-806, 1996. Denatured RNA was size
                   fractionated on a 1% agarose gel. First strand cDNA
                   synthesis was primed with oligo-dT primer containing a Not
                   I site. Double strand cDNA was size selected according to
                   mRNA size fraction, ligated with EcoR I adaptor, digested
                   with NotI and then cloned directionally into pYX-Asc
                   vector. The library tag sequence located between the Not I
                   site and the polyA tail is CATCTCTACT. This library was
                   created for the University of Iowa Program for Rat Gene
                   Discovery and Mapping (Val Sheffield, Bento Soares and Tom
                   Casavant)."
                       168 c
                               154 g
                                        119 t
                                                  2 others
BASE COUNT
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                        81.4%; Pred. No. 1.9e-74;
  Best Local Similarity
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                                                             0; Gaps
                                                                        0;
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Qy
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Db
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Qу
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Db
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Db
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Qу
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Db
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Qу
            \perp 11
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Db
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Qу
                             ł
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LOCUS
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          AI119218
ACCESSION
          AI119218.1 GI:3519542
VERSION
KEYWORDS
          EST.
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SOURCE
 ORGANISM Mus musculus
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          Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
             (bases 1 to 816)
REFERENCE
          Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
 AUTHORS
          Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
          Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,
          Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and
          Waterston, R.
          The WashU-HHMI Mouse EST Project
  TITLE
          Unpublished
  JOURNAL
          Contact: Marra M/Mouse EST Project
COMMENT
          WashU-HHMI Mouse EST Project
          Washington University School of MedicineP
          4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
          Tel: 314 286 1800
          Fax: 314 286 1810
          Email: mouseest@watson.wustl.edu
          This clone is available royalty-free through LLNL; contact the
          IMAGE Consortium (info@image.llnl.gov) for further information.
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MGI:936407
Seq primer: custom primer used
High quality sequence stop: 473.

FEATURES
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Site\_2: DraIII (CACCATGTG); 1st strand cDNA was primed with an oligo(dT) primer [ATGTGGCCTTTTTTTTTTTTTTT];
double-stranded cDNA was ligated to a DraIII adaptor [TGTTGGCCTACTGG], digested and cloned into distinct DraIII sites of the pME18S-FL3 vector (5' site CACTGTGTG, 3' site CACCATGTG). XhoI should be used to isolate the cDNA insert. Size selection was performed to exclude fragments <1.5kb. Library constructed by Dr. Sumio Sugano (University of Tokyo Institute of Medical Science).

0;

(University of Tokyo Institute of Medical Science Custom primers for sequencing: 5' end primer CTTCTGCTCTAAAAGCTGCG and 3' end primer

CGACCTGCAGCTCGAGCACA."

BASE COUNT ORIGIN

230 a 219 c 172 g 187 t 8 others

Query Match 64.9%; Score 339.2; DB 9; Length 816; Best Local Similarity 81.2%; Pred. No. 2.5e-74; Matches 389; Conservative 0; Mismatches 90; Indels 0; Gaps

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Qу
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Db
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Db
RESULT 7
BF383724
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                                                             EST 27-NOV-2000
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DEFINITION
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ACCESSION
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           BF383724.1 GI:11365029
VERSION
KEYWORDS
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SOURCE
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 ORGANISM
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           1 (bases 1 to 594)
REFERENCE
           NIH-MGC http://mgc.nci.nih.gov/.
 AUTHORS
           National Institutes of Health, Mammalian Gene Collection (MGC)
 TITLE
           Unpublished
 JOURNAL
           Contact: Robert Strausberg, Ph.D.
COMMENT
           Email: cgapbs-r@mail.nih.gov
           Tissue Procurement: Jeffrey E. Green, M.D.
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
           found through the I.M.A.G.E. Consortium/LLNL at:
           http://image.llnl.gov
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           High quality sequence stop: 589.
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                    Technologies. Note: this is a NCI CGAP Library."
BASE COUNT
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Qу	136	GAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTACTGTGCACCCCTCAAG 195					
Db	227	GAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGACTGGAGATGTACTGTGCCCCACTGAAG 286					
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Db	287						
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Db	407						
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Qу	436	TGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTTCAAAGATGG 489					
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Qу	490	CATTTCCC 497					
Db	587						
RESULT 8 CB959991 LOCUS DEFINITION ACCESSION VERSION KEYWORDS SOURCE ORGANIS REFERENCE AUTHORS TITLE JOURNAL COMMENT	M H H I I I I I I I I I I I I I I I I I	796 bp mRNA linear EST 29-APR-2003 MGENCOURT_13888044 NIH_MGC_147 Homo sapiens cDNA clone MAGE:30341081 5', mRNA sequence. MS959991 MS959991.1 GI:30216107 MST. Momo sapiens (human) Momo sapiens Mukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  (bases 1 to 796) MIH-MGC http://mgc.nci.nih.gov/. Mational Institutes of Health, Mammalian Gene Collection (MGC) Mpublished Montact: Robert Strausberg, Ph.D. Mail: cgapbs-r@mail.nih.gov Missue Procurement: Dr. Stefan Hansson CDNA Library Preparation: Michael J. Brownstein (NHGRI) with help					
	ā	and advice from Piero Carninci (RIKEN)					

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Agencourt Bioscience Corporation Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Plate: NDAM371 row: p column: 18 High quality sequence stop: 707. Location/Qualifiers FEATURES 1. .796 source /organism="Homo sapiens" /mol type="mRNA" /db xref="taxon:9606" /clone="IMAGE:30341081" /tissue\_type="Human Placenta" /lab host="DH10B TonA" /clone lib="NIH MGC 147" /note="Organ: placenta; Vector: pBluescriptR; Site 1: alI-XhoI; Site 2: BamH; Oligo-dT primed using primer 5'-TTTTTTTTTTTTTTTTTVN-3', size-selected for average insert size 2.3 kb and normalized to ROT 5. This is a primary library enriched for full-length clones and constructed using the Cap-trapper method (Carninci, in preparation). Library constructed by M. Brownstein (NIMH/NHGRI, National Institutes of Health). Note: This is a NIH MGC library." 197 c 191 g 184 t 224 a BASE COUNT ORIGIN 63.9%; Score 334.4; DB 14; Length 796; Query Match 84.6%; Pred. No. 3.9e-73; Best Local Similarity 0; Mismatches 26; Indels 55: Gaps 4; Matches 445; Conservative 1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGAGACCCTCTTCAGTTCGTGTGTGGAGAC 60 Qу 180 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 239 Db 61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120 Qу 240 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 299 Db 121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180 Qу 300 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 359 Db Qу 360 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 419 Db 241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300 Qу 11111111111 111 420 ATGCCCAAGACCCAG----- 434 Db 301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360 Qy 435 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 487 Db

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Qy

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              (bases 1 to 499)
REFERENCE
           Bonaldo, M.F., Lennon, G. and Soares, M.B.
 AUTHORS
           Normalization and subtraction: two approaches to facilitate gene
 TITLE
           Genome Res. 6 (9), 791-806 (1996)
 JOURNAL
           97044477
 MEDLINE
           8889548
  PUBMED
COMMENT
           Contact: Chin, H
           National Institute of Mental Health
           6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD
           20892-9643, USA
           Tel: 301 443 1706
           Fax: 301 443 9890
           Email: mEST@mail.nih.gov
           The sequence contained an oligo-dT track that was present in the
           oligonucleotide that was used to prime the synthesis of first
           strand cDNA and therefore this may represent a bonafide poly A
           tail. The sequence tag present in the cDNA between the NotI site
           and the oligo-dT track served to identify it as a clone from the
           normalized pineal glands library cDNA Library Preparation: M.B.
           Soares Lab Clone distribution: Researchers may obtain BMAP cDNA
           clones from RESEARCH GENETICS. It should be noted that Bento Soares
           is generating a small number of additional specialized
           non-redundant arrays of BMAP cDNAs whose availability will be
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           Seg primer: M13 Forward
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                 ultimately derived from a mixture of individually tagged
                 normalized libraries from ten regions of the mouse brain
                 (cerebellum, brain stems, olfactory bulbs, hypothalamus,
                 cortex, amygdala, basal ganglia, pineal gland, striatum,
                 hipoccampus) after a series of subtractions to reduce the
                 representation of cDNAs from which ESTs had already been
                 generated. The following serially subtracted libraries
                 were generated in this process: NIH BMAP M S4,
                 NIH BMAP M S3.3, NIH BMAP M S3.2, NIH BMAP M S3.1,
                 NIH BMAP M S2, NIH BMAP M S1. The subtracted library
                 (NIH BMAP M S4) was constructed as follows: PCRamplified
                 cDNA inserts from NIH_BMAP M S3.3, NIH BMAP M S3.2, and
                 NIH BMAP M S3.1 clones from which 3' ESTs had been derived
                 was used as a driver in a hybridization with a pool of
                 the NIH BMAP M S3.3, NIH BMAP M S3.2, and NIH BMAP M S3.1
                 libraries in the form of single-stranded circles. The
                 remaining single-stranded circles (subtracted library)
                 was purified by hydroxyapatite column chromatography,
                 converted to double-stranded circles and electroporated
                 into DH10B bacteria (LifeTechnologies) to generate the
                 NIH BMAP M S4 library. This procedure has been previously
                 described (Bonaldo, Lennon and Soares, Genome Research
                 6:791-806, 1996)
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                 TAG SEQ=CAGAC"
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Best Local Similarity
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Matches 396; Conservative
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                                                Indels
                                                          7; Gaps
                                                                     2;
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                     499 TGTGTGGACCGAGGGGCTTTTACTTCAACAAGCCCACAGGCTATGGCTCCAGCATTCGGA 440
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       170 TGGAGATGTACTGTGCACCCCTCAAGCCGGCAAAGGCAGCCCGCTCCGTCCCTGCCCAGC 229
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230 GCCACACCGACATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGA 289

290 AGTCTCAGAGGAGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGA 349

BASE COUNT

Query Match

ORIGIN

Qy

Db

Qу

Db

Qу

Db

Qу

Db

Qу

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259 AGCTGCAAAGGAGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGAAGTGCAGGA 200
Db
         350 AACAAGAACTACAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGC 409
Qу
             1 11 1 11
         199 AACAAGACCTACAGAATGTAGGAGGAGCCTCCCACGGAGCAGAAAATGCCACATCACCGC 140
Db
         410 AGGACCCTTTGCTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA-----AAAAA 463
Qу
                                        1111 1111111
                                   1
         139 AGGATCCTTTGCTGCTTGAGCAACCTGCAAAACATCGAAACACCTACCAAATAACAATAA 80
Db
         464 TAAGTTTGATCACATTTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATT 522
Qу
                    79 TAAGTCCAATAACATTACAAAGATGGGCATTTCCCCCAATGAAATATACAAGTAAACATT 20
Db
         523 C 523
Qу
          19 C 19
Db
RESULT 10
AI876493/c
                                  642 bp
                                            mRNA
                                                   linear
                                                            EST 21-JUL-1999
LOCUS
           AI876493
          uj59b10.x1 Sugano mouse liver mlia Mus musculus cDNA clone
DEFINITION
           IMAGE: 1924219 3' similar to gb: X57025 rna1 INSULIN-LIKE GROWTH
           FACTOR IA PRECURSOR (HUMAN); gb:X04482 Mouse mRNA for
           preproinsulin-like growth factor IB (MOUSE);, mRNA sequence.
ACCESSION
           AI876493
           AI876493.1 GI:5550542
VERSION
           EST.
KEYWORDS
           Mus musculus (house mouse)
SOURCE
 ORGANISM Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
           1 (bases 1 to 642)
           Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T.,
 AUTHORS
           Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person
           ,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter
           ,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
           Waterston, R. and Wilson, R.
           The WashU-NCI Mouse EST Project 1999
 TITLE
 JOURNAL
           Unpublished
           Contact: Marra M/WashU-NCI Mouse EST Project 1999
COMMENT
           Washington University School of Medicine
           4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
           Tel: 314 286 1800
           Fax: 314 286 1810
           Email: mouseest@watson.wustl.edu
           This clone is available royalty-free through LLNL; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
           MGI:980511
           Seq primer: custom primer used
           High quality sequence stop: 257.
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                   /mol type="mRNA"
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                /clone="IMAGE:1924219"
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                /dev stage="adult"
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                /clone lib="Sugano mouse liver mlia"
                /note="Organ: liver; Vector: pME18S-FL3; Site_1: DraIII
                 (CACTGTGTG); Site 2: DraIII (CACCATGTG); 1st strand cDNA
                was primed with an oligo(dT) primer
                 [ATGTGGCCTTTTTTTTTTTTTTTT]; double-stranded cDNA was
                ligated to a DraIII adaptor [TGTTGGCCTACTGG], digested
                and cloned into distinct DraIII sites of the pME18S-FL3
                vector (5' site CACTGTGTG, 3' site CACCATGTG). XhoI should
                be used to isolate the cDNA insert. Size selection was
                performed to exclude fragments <1.5kb. Library
                constructed by Dr. Sumio Sugano (University of Tokyo
                Institute of Medical Science). Custom primers for
                sequencing: 5' end primer CTTCTGCTCTAAAAGCTGCG and 3' end
                primer CGACCTGCAGCTCGAGCACA."
BASE COUNT
            127 a
                    154 c
                           175 q
                                  185 t
                                           1 others
ORIGIN
                    61.3%;
                           Score 320.8; DB 9; Length 642;
 Query Match
 Best Local Similarity
                    80.1%;
                           Pred. No. 9.6e-70;
 Matches 403; Conservative
                          0; Mismatches
                                        93:
                                            Indels
                                                       Gaps
                                                              2;
         2 GACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGACA 61
Qy
           503 GACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGATGCTCTTCAGGTCGTGTGTGGACCGA 444
Db
        62 GGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAGA 121
Qу
                   Db
        443 GGGGCTTTTTCTTCAACAAGGCCACAGGCTATGGCTCCAGCATTTGGAGGGCACCTCAGA 384
        122 CAGGCATCGTGGATGAGTGCTGCTTCCGG-AGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
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           383 CAGTCAATGTGGATGAGTGTTGCTTCCGGAAGCTGTGATCTGAGAAGACTGNAGATGTAC 324
Db
        Qy
           323 TGTGCCCCACTGAAGCCTACAAAAGCAGCCCGCTCTATCCGTGCCCAGCGCCACACTGAC 264
Db
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Qу
           Db
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Qy
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361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420

143 CAGAATGTAGGAGGAGCCTCCCACGGAGCAGAAAATGCCACATCACCGCAGGATCCTTTG 84

421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA-----AAAAATAAGTTTGATC 474

1

Db

Qу

Db

Qу

 $\mathbf{I}$ 

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83 CTGCTTGAGCAACCTGCAAAACATCGAAACACCTACCAAATAACAATAATAAGTCCAATA 24
Db
          475 ACATTTCAAAGATGGCATTTCCC 497
Qу
              11 11
           23 ACATTACAAAGATGGGCATTTCC 1
Db
RESULT 11
BM984670/c
                                                                EST 20-FEB-2003
                                                       linear
                                     673 bp
                                               mRNA
LOCUS
            BM984670
           UI-CF-EC1-abj-k-24-0-UI.sl UI-CF-EC1 Homo sapiens cDNA clone
DEFINITION
            UI-CF-EC1-abj-k-24-0-UI 3', mRNA sequence.
            BM984670
ACCESSION
            BM984670.1 GI:19610417
VERSION
KEYWORDS
            EST.
            Homo sapiens (human)
SOURCE
  ORGANISM Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
            1 (bases 1 to 673)
            Bonaldo, M.F., Lennon, G. and Soares, M.B.
  AUTHORS
            Normalization and subtraction: two approaches to facilitate gene
  TITLE
            discovery
            Genome Res. 6 (9), 791-806 (1996)
  JOURNAL
            97044477
  MEDLINE
            8889548
   PUBMED
            Contact: McCray, PB
COMMENT
            McCray Lab
            University of Iowa
            2024 University of Iowa Med Labs, Iowa City, IA 52242, USA
            Tel: 319 356 4866
            Fax: 319 356 7171
            Email: paul-mccray@uiowa.edu
            Tissue Procurement: Dr. M. J. Welsh, University of Iowa
             cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
             cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
             DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
             Clone Distribution: Researchers may obtain clones from Research
            Genetics (www.resgen.com) or from Open Biosystems
            (www.openbiosystems.com).
            Seg primer: M13 FORWARD
            POLYA=Yes.
                     Location/Oualifiers
FEATURES
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                     /tissue type="Lung"
                     /dev stage="Adult and Fetal"
                     /lab host="DH10B (Life Technologies) (T1 phage resistant)"
                     /clone lib="UI-CF-EC1"
                     /note="Organ: Lung; Vector: pT7T3-Pac (Pharmacia) with a
                     modified polylinker; Site_1: EcoR I; Site_2: Not I;
                     UI-CF-EC1 is a normalized cDNA library containing the
                     following tissue(s): Normal lung from adult and from fetal
                     day 64, day 87, week 19 and week 42. The library was
```

constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pT7T3-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is AAGTGCTTAC.

TAG LIB=UI-CF-EC1

TAG\_TISSUE=Normal Lung Epithelial Cells Tissue nos 369-371 and 380-383

TAG SEQ=AAGTGCTTAC"

BASE COUNT

152 a 164 c 169 g 188 t

ORIGIN

61.3%; Score 320.8; DB 12; Length 673; Query Match Best Local Similarity 84.2%; Pred. No. 9.7e-70; Matches 443; Conservative 0; Mismatches 27; Indels 56; Gaps 5; 1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60 Qy 492 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 433 Db 61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120 Qу 432 AGGGG-TTTTATTTCAGCAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 374 Db 121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180 Qy 373 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 314 Db Qу 313 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 254 Db 241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300 Qу 111111111111111 253 ATGCCCAAGACCCAG----- 239 Db 301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360 Qу 238 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 186 Db 361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420 Qу 185 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 126 Db 421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478 Qу 125 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 66 Db 479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523 Qу 65 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 20 Db

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RESULT 12
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                                                             EST 01-DEC-1998
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                                                     linear
                                   575 bp
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LOCUS
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DEFINITION
           clone IMAGE: 1849953 3' similar to gb: X57025 rnal INSULIN-LIKE
           GROWTH FACTOR IA PRECURSOR (HUMAN);, mRNA sequence.
           AI248089
ACCESSION
           AI248089.1 GI:3843486
VERSION
           EST.
KEYWORDS
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SOURCE
           Homo sapiens
  ORGANISM
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           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
              (bases 1 to 575)
           NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS
           National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 TITLE
           Tumor Gene Index
  JOURNAL
           Unpublished
           Contact: Robert Strausberg, Ph.D.
COMMENT
           Email: cgapbs-r@mail.nih.gov
           This clone is available royalty-free through LLNL ; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
                                Std Error: 0.00
           Insert Length: 918
           Seq primer: -40UP from Gibco
           High quality sequence stop: 380.
                    Location/Qualifiers
FEATURES
                    1. .575
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                    This is a subtracted version of the original Soares fetal
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                    liver spleen 1NFLS library.
                    with a Pac I - oligo(dT) primer [5'
                    double-stranded cDNA was ligated to Eco RI adaptors
                    (Pharmacia), digested with Pac I and cloned into the Pac I
                    and Eco RI sites of the modified pT7T3 vector. Library
                    went through one round of normalization. Library
                    constructed by Bento Soares and M. Fatima Bonaldo."
                                         156 t
                                                    1 others
BASE COUNT
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                                 131 g
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                                                              55;
                                                                          4;
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                                                28;
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Qу
             551 TGCGGGGCTGAGCTGGTGNATGCTCTTCAGTTCGTGTGAAGACAGGGGCTTTTATTTC 492
Db
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Qy	76 AACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAGACAGGCATCGTGGAT 135					
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Qу	136 GAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTACTGTGCACCCCTCAAG 195					
Db	431 GAGTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTATTGCGCACCCCTCAAG 372					
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Db	303 ACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTACAGGATGTAGGAAGA 244					
Qу	376 CCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTGCTCTGCAC-AGTTAC 434					
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Qy	435 CTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTTCAAAGAT-GGCAT 492					
Db	183 CTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACATTTAAAAGATGGGCGT 124					
Qy	493 TTCCCCCAATGAAATACACAAGTAAACATTC 523					
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AUTHORS	Lee, N.H., Glodek, A., Chandra, I., Mason, T.M., Quackenbush, J.,					
TITLE	<pre>Kerlavage,A.R. and Adams,M.D. Rat Genome Project: Generation of a Rat EST (REST) Catalog &amp; Rat Gene Index</pre>					
JOURNAL	npublished					
COMMENT	Other_ESTs: TC50779 Contact: Lee, NH					
	The Institute for Genomic Research 9712, Medical Center Drive, Rockville, MD 20850, USA					

```
Tel: (301)-838-3529
         Fax: (301)-838-0208
         Email: nhlee@tigr.org
         Seq primer: M13-21.
                Location/Qualifiers
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                Site 2: NotI"
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 Query Match
 Best Local Similarity 81.8%; Pred. No. 7.2e-67;
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 Matches 383; Conservative
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           468 GGGCTTTTACTTCAACAAGCCCACAGGCTATGGCTCCAGCATTCGGAGGGCACCACAGAC 409
Db
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Qу
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Qу
           168 GAATGTAGGAGGAGCCTCCCGAGGAACAGAAAATGCCACGTCACCGCAAGATCCTTTGCT 109
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Qу
                       108 GCTTGAGCAACCTGCAAAACATCGGAACACCTGCCAAATATCAATAATGAGTTCAATACC 49
Db
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Db
RESULT 14
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AA542914/c

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            (HUMAN);, mRNA sequence.
ACCESSION
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VERSION
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SOURCE
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            1 (bases 1 to 498)
REFERENCE
           NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS
           National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 TITLE
            Tumor Gene Index
            Unpublished
  JOURNAL
            Contact: Robert Strausberg, Ph.D.
COMMENT
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Michael J. Brownstein, M.D., Ph.D., Michael R.
            Emmert-Buck, M.D., Ph.D.
             cDNA Library Preparation: M. Bento Soares, Ph.D.
             cDNA Library Arrayed by: Greg Lennon, Ph.D.
             DNA Sequencing by: Washington University Genome Sequencing Center
             Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www-bio.llnl.gov/bbrp/image/image.html
            Insert Length: 603
                               Std Error: 0.00
            Seq primer: -40ml3 fwd. ET from Amersham
            High quality sequence stop: 412.
                    Location/Qualifiers
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                     /lab host="DH10B"
                     /clone lib="NCI CGAP Pr21"
                     /note="Organ: prostate; Vector: pT7T3D-Pac (Pharmacia)
                     with a modified polylinker; 1st strand cDNA was prepared
                     from normal prostate bulk tissue, and was then primed with
                     a Not I - oligo(dT) primer. Double-stranded cDNA was
                     ligated to Eco RI adaptors (Pharmacia), digested with Not
                     I and cloned into the Not I and Eco RI sites of the
                     modified pT7T3 vector. Library is not normalized. Library
                     was constructed by Bento Soares and M. Fatima Bonaldo. "
BASE COUNT
                105 a
                         135 с
                                  123 q
                                           135 t
ORIGIN
  Query Match
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  Best Local Similarity
                         83.5%; Pred. No. 8.2e-66;
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Qу
                          476 GGACCGGAGAACTTTTGCGGGGCTTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGA 417
Db
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0	<b>C</b> 0	CAGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCA 119
ДУ		CAGGGGCTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCGCTGT 113
Db		
QУ	120	GACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTA 179
Db	357	GACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTA 298
QУ	180	CTGTGCACCCCTCAAGCCGGCAAAGGCAGCCCGCTCCGTCCG
Db	297	TTGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGA 238
Qу	240	CATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAG 299
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Db	221	AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACT 170
Qy	360	ACAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTT 419
Db	169	ACAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTT 110
Qу	420	GCTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACA 477
Db	109	GCTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACA 50
Qу	478	TTTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Db	49	TTTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 3
RESULT 15 AI604642 LOCUS DEFINITIO	N V C F	1604642 882 bp mRNA linear EST 21-APR-1999 m43d08.yl Stratagene mouse diaphragm (#937303) Mus musculus cDNA lone IMAGE:1001007 5' similar to gb:M11568 INSULIN-LIKE GROWTH ACTOR IB PRECURSOR (HUMAN); gb:X04482 Mouse mRNA for
ACCESSION	_	reproinsulin-like growth factor IB (MOUSE);, mRNA sequence. I604642
VERSION		I604642.1 GI:4613809
KEYWORDS SOURCE		ST. us musculus (house mouse)
ORGANIS	M M	us musculus ukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
	М	ammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE AUTHORS	M U	(bases 1 to 882)  Garra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T., Inderwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk, R., Ritter E., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R.,  Gaterston, R. and Wilson, R.
TITLE	Т	he WashU-NCI Mouse EST Project 1999
JOURNAL COMMENT	C	npublished Contact: Marra M/WashU-NCI Mouse EST Project 1999 Cashington University School of Medicine
	-	

```
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
          Tel: 314 286 1800
          Fax: 314 286 1810
          Email: mouseest@watson.wustl.edu
          This clone is available royalty-free through LLNL; contact the
          IMAGE Consortium (info@image.llnl.gov) for further information.
          MGI:565223
          This read is a RESEQUENCE of a previously sequenced mouse clone
          This read has been verified (found to hit its original self in the
          correct orientation)
          Seq primer: -40RP from Gibco
          High quality sequence stop: 361.
                 Location/Oualifiers
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Search completed: December 13, 2003, 07:29:51

Job time : 1704.81 secs

## GenCore version 5.1.6 Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

December 13, 2003, 05:41:20; Search time 2336.77 Seconds Run on:

(without alignments)

9156.102 Million cell updates/sec

US-09-852-261-5 Title:

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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5: gb ov:\*

gb pat:\*

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gb\_pr:\* 9:

10: gb\_ro:\*

11: gb sts:\*

12: gb\_sy:\*

13: gb un:\* 14: gb vi:\*

15: em ba:\*

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18: em in:\*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

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2	523	100.0	523	6	AX300783	AX300783 Sequence
3	467.4	89.4	517	6	AX147742	AX147742 Sequence
4	467.4	89.4	517	6	AX300779	AX300779 Sequence
5	409	78.2	471	6	AX147754	AX147754 Sequence
6	409	78.2	471	6	AX300791	AX300791 Sequence
7	364.8	69.8	1536	10	BC012409	BC012409 Mus muscu
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ACCESSION
            AX147746
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KEYWORDS
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REFERENCE
  AUTHORS
            Goldspink, G.R. and Johnson, I.R.
            Use of the insulin-like-growth factor i isoform mgf for the
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REFERENCE
         Goldspink, G.D. and Terenghi, G.B.
 AUTHORS
         Repair of nerve damage
 TITLE
         Patent: WO 0185781-A 5 15-NOV-2001;
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REFERENCE
         Goldspink, G.R. and Johnson, I.R.
 AUTHORS
 TITLE
         Use of the insulin-like-growth factor i isoform mgf for the
         treatment of neurological disorders
         Patent: WO 0136483-A 1 25-MAY-2001;
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REFERENCE
          Goldspink, G.D. and Terenghi, G.B.
 AUTHORS
          Repair of nerve damage
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REFERENCE
 AUTHORS
         Goldspink, G.R. and Johnson, I.R.
         Use of the insulin-like-growth factor i isoform mgf for the
 TITLE
         treatment of neurological disorders
         Patent: WO 0136483-A 13 25-MAY-2001;
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Qу	1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60	
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

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Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
REFERENCE
 AUTHORS
        Goldspink, G.D. and Terenghi, G.B.
 TITLE
        Repair of nerve damage
 JOURNAL,
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DEFINITION
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            MGC:18617 IMAGE:4194295), complete cds.
            BC012409
ACCESSION
            BC012409.1 GI:15214568
VERSION
KEYWORDS
            MGC.
            Mus musculus (house mouse)
SOURCE
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REFERENCE
               (bases 1 to 1536)
 AUTHORS
            Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
            Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D.,
            Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,
            Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,
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            Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S.,
            Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J.,
            Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J.,
            McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S.,
            Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W.,
            Villalon, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A.,
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            Butterfield, Y.S., Krzywinski, M.I., Skalska, U., Smailus, D.E.,
            Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.
            Generation and initial analysis of more than 15,000 full-length
 TITLE
            human and mouse cDNA sequences
            Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
  JOURNAL
 MEDLINE
            22388257
            12477932
   PUBMED
            2 (bases 1 to 1536)
REFERENCE
            Strausberg, R.
 AUTHORS
  TITLE
            Direct Submission
            Submitted (15-AUG-2001) National Institutes of Health, Mammalian
  JOURNAL
            Gene Collection (MGC), Cancer Genomics Office, National Cancer
            Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
            USA
            NIH-MGC Project URL: http://mgc.nci.nih.gov
  REMARK
            Contact: MGC help desk
COMMENT
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Jeffrey E. Green, M.D.
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Baylor College of Medicine Human Genome
```

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Center code: BCM-HGSC
          Web site: http://www.hgsc.bcm.tmc.edu/cdna/
          Contact: amg@bcm.tmc.edu
          Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Loulseged, H.,
          Kowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Nanavati,
          A.N., Gibbs, R.A.
          Clone distribution: MGC clone distribution information can be found
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Sequencing Center

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ACCESSION
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          X06108.1 GI:56426
VERSION
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SOURCE
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          Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
          Rattus.
REFERENCE
          1
          Shimatsu, A. and Rotwein, P.
 AUTHORS
          Sequence of Two Rat Insulin-like Growth Factor I mRNAs Differing
 TITLE
          Within the 5' Untranslated Region
          Nucleic Acids Res. 15 (1987) In press
 JOURNAL
            (bases 1 to 798)
REFERENCE
          2
 AUTHORS
          Rotwein, P.
 TITLE
          Direct Submission
          Submitted (21-OCT-1987) Rotwein P., Washington University, School
 JOURNAL
          of Medicine, 660 South Euclid Avenue, Box 8127, St. Louis, MO
          63110. USA
          Another IGF-I mRNA of rat liver differing in the 5' UT-region is
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           X06107 M32260 Y00429
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VERSION
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REFERENCE
            1
  AUTHORS
           Shimatsu, A. and Rotwein, P.
            Sequence of two rat insulin-like growth factor I mRNAs differing
  TITLE
           within the 5' untranslated region
           Nucleic Acids Res. 15 (17), 7196 (1987)
  JOURNAL
           88015572
  MEDLINE
           3658684
   PUBMED
              (bases 1 to 958)
REFERENCE
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           Rotwein, P.
  AUTHORS
           Direct Submission
  TITLE
            Submitted (21-OCT-1987) Rotwein P., Washington University, School
  JOURNAL
            of Medicine, 660 South Euclid Avenue, Box 8127, St. Louis, MO
            63110, USA
COMMENT
            Another IGF-I mRNA of rat liver differing in the 5' UT-region is
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JOURNAL MEDLINE PUBMED	LeRoith,D.  Molecular cloning of rat insulin-like growth factor I complementary deoxyribonucleic acids: differential messenger ribonucleic acid processing and regulation by growth hormone in extrahepatic tissues Mol. Endocrinol. 1 (3), 243-248 (1987). 88288198 3453891
COMMENT	Original source text: Rat (Sprague-Dawley) adult liver cDNA to mRNA, clone pRIGF-1-42. Draft entry and computer-readable copy of sequence in [Mol. Endocrinol. (1987) In press] kindly

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provided by S.R.Lasky, 16-MAR-1987.
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DEFINITION
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ACCESSION
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VERSION
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 ORGANISM
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           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
           Rattus.
REFERENCE
           1
 AUTHORS
           Goldspink, G.R. and Johnson, I.R.
           Use of the insulin-like-growth factor i isoform mgf for the
 TITLE
           treatment of neurological disorders
           Patent: WO 0136483-A 3 25-MAY-2001;
 JOURNAL
           University College London (GB)
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REFERENCE
         Goldspink, G.D. and Terenghi, G.B.
 AUTHORS
         Repair of nerve damage
 TITLE
         Patent: WO 0185781-A 3 15-NOV-2001;
 JOURNAL
         University College London (GB); East Grinstead Medical Research
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#### RESULT 13 MMIGFIBR

Qу

Db

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DEFINITION Mouse mRNA for preproinsulin-like growth factor IB.

ACCESSION X04482

VERSION X04482.1 GI:51806

KEYWORDS growth factor; insulin-like growth factor IB; preproinsulin-like

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growth factor IB; signal peptide.
SOURCE
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  ORGANISM
           Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
              (bases 1 to 651)
  AUTHORS
            Bell, G.I., Stempien, M.M., Fong, N.M. and Rall, L.B.
  TITLE
            Sequences of liver cDNAs encoding two different mouse insulin-like
            growth factor I precursors
  JOURNAL
           Nucleic Acids Res. 14 (20), 7873-7882 (1986)
  MEDLINE
            87040760
   PUBMED
            3774549
           The sequence is identical to the preproIGF-IA sequence (X04480)
COMMENT
            except for the presence of a 52 bp insertion following codon 86
            (position 397 to 448), caused by alternative RNA splicing. The B
            domain of IGF comprises residues 1-29 (position 139-225), the C
            domain residues 30-41 (position 226-261), the A domain residues
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REFERENCE
 AUTHORS
         Baak, J. and Mutter, G.L.
 TITLE
         Prognostic classification of breast cancer
 JOURNAL
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#### GenCore version 5.1.6 Copyright (c) 1993 - 2003 Compugen Ltd.

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5	467.4	89.4	517	24	AAS16877	Human mechano-grow
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7	409	78.2	471	24	AAS16884	Rabbit insulin-lik
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45	193	36.9	210	24	ABA03146	Native mature IGF-

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RESULT 1
AAD06400
     AAD06400 standard; cDNA; 523 BP.
TD
XX
AC
     AAD06400;
XX
DΤ
     10-AUG-2001 (first entry)
XX
DΕ
     Rabbit IGF-I isoform mechano-growth factor (MGF) cDNA.
XX
ΚW
     Rabbit; IGF-I isoform; Insulin-like Growth Factor-I; MGF;
     mechano-growth factor; neurological disorder; neurodegenerative disorder;
KW
     amyotrophic lateral sclerosis; spinal muscular atrophy; muscular atrophy;
KW
     poliomyelitis; post-polio syndrome; toxin; motoneurone disorder;
KW
     nerve damage; autosomal muscular dystrophy; diabetic neuropathy;
KW
     sex-linked muscular dystrophy; peripheral neuropathy;
KW
     Alzheimer's disease; Parkinson's disease; ss.
KW
XX
OS
    Oryctolagus cuniculus.
XX
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     CDS
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                     /product= "Mechano-growth factor (MGF)"
FT
FT
                     /note= "This region comprises exons 3-6. The CDS does
FT
                     not include start codon"
FT
                     /partial
XX
ΡN
    WO200136483-A1.
XX
     25-MAY-2001.
PD
XX
PF
    15-NOV-2000; 2000WO-GB04354.
XX
PR
     15-NOV-1999;
                    99GB-0026968.
XX
     (UNLO ) UNIV COLLEGE LONDON.
PA
XX
PI
    Goldspink G, Johnson I;
XX
DR
    WPI; 2001-355620/37.
    P-PSDB; AAE02449.
DR
XX
     Use of mechano-growth factor, an isoform of Insulin-like Growth
PT
     Factor-I, capable of reducing motoneurone loss, in the manufacture of a
PT
    medicament for the treatment of neurological disorder -
PT
XX
    Claim 4; Page 53-54; 66pp; English.
PS
XX
    The present invention relates to use of mechano-growth factor (MGF),
CC
CC
     an Insulin-like Growth Factor-I (IGF-I) isoform in the manufacture of a
CC
    medicament for the treatment of neurological disorder. The MGF is capable
CC
     of reducing motoneurone loss by 20% or greater in response to nerve
CC
     avulsion, and effects motoneurone rescue, preferably adult motoneurone
CC
     rescue. The MGF polynucleotide and polypeptide are useful in the
CC
    manufacture of a medicament for the treatment of a neurological disorder,
     including a disorder of motoneurones and/or neurodegenerative disorder,
CC
```

```
CC
    e.g., amyotrophic lateral sclerosis, spinal muscular atrophy, progressive
CC
    spinal muscular atrophy, infantile or juvenile muscular atrophy,
    poliomyelitis or post-polio syndrome, a disorder caused by exposure to a
CC
    toxin, motoneurone trauma, a motoneurone lesion or nerve damage, an
CC
    injury that affects motoneurones, motoneurone loss associated with aging,
CC
CC
    autosomal or sex-linked muscular dystrophy, diabetic neuropathy,
CC
    peripheral neuropathies, Alzheimer's disease and Parkinson's disease.
    The present sequence is rabbit IGF-I isoform MGF cDNA. MGF is a muscle
CC
CC
    isoform having extracellular (Ec) domain, hence also referred as
    IGF-I-Ec. The MGF protein comprises amino acid sequences encoded by
CC
CC
    nucleic acid sequence of IGF-I exons 4, 5 and 6 in the reading frame
    of MGF.
CC
XX
    Sequence 523 BP; 154 A; 129 C; 142 G; 98 T; 0 other;
SO
 Query Match
                    100.0%; Score 523; DB 22;
                                          Length 523;
 Best Local Similarity
                    100.0%; Pred. No. 5.1e-144;
 Matches 523; Conservative
                         0: Mismatches
                                       0: Indels
                                                     Gaps
                                                           0;
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           1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Db
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Qу
           Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
       121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
       QУ
          Db
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Qу
           241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
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Qу
          301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
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Qу
          361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Db
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Qу
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Db
       481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Db
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     AAS16879 standard; cDNA; 523 BP.
XX
     AAS16879;
AC
XX
DT
     25-FEB-2002 (first entry)
XX
DΕ
     Rabbit mechano-growth factor (MGF) cDNA.
XX
ΚW
     Rabbit; mechano-growth factor; insulin-like growth factor I; IGF-I; MGF;
KW
     neuroprotective; nerve damage; peripheral nervous system; nerve severing;
     muscle; neurological disorder; motoneuron loss; motorneuron disorder; ss;
KW
     nerve avulsion.
KW
XX
OS
     Oryctolagus cuniculus.
XX
FΗ
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     Key
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     CDS
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                     /product= "Rabbit MGF"
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                     /partial
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FT
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FT
FT
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FT
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FT
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PN
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XX
PD
     15-NOV-2001.
XX
PF
     10-MAY-2001; 2001WO-GB02054.
XX
     10-MAY-2000; 2000GB-0011278.
PR
XX
PΑ
     (UNLO ) UNIV COLLEGE LONDON.
PΑ
     (EGRI-) EAST GRINSTEAD MEDICAL RES TRUST.
XX
PΙ
     Goldspink G, Terenghi G;
XX
DR
     WPI; 2002-055585/07.
     P-PSDB; AAU10561.
DR
XX
PT
     Use of insulin-like growth factor I (IGF-I) isoform known as
     mechano-growth factor which is encoded by IGF-I exons 4,5,6 and has
PT
PT
     ability to reduce motoneuron loss in response to nerve avulsion, to
PT
     treat nerve damage -
XX
PS
     Disclosure; Fig 7; 65pp; English.
```

```
XX
CC
    The invention relates to the use of an insulin-like growth factor I
CC
    (IGF-I) isoform, known as mechano-growth factor (MGF), in the manufacture
CC
    of a medicament for treating nerve damage in the peripheral nervous
CC
    system, or for treating nerve damage by localising MGF at the site of
    damage. The nerve damage may include severing of a nerve. The treatment
CC
CC
    may be combined with another treatment (such as a polypeptide growth
    factor other than MGF) that prevents or diminishes degeneration of the
CC
    target organ (for example, muscle) which the damaged nerve innervates,
CC
    whereby the treatment of the muscle with MGF or a polynucleotide encoding
CC
CC
    MGF prevents or diminishes degeneration. The method is useful for
CC
    treating neurological disorders, preferably motorneuron disorders. These
    methods can reduce motoneuron loss by 20% or greater in response to nerve
CC
CC
    avulsion. This sequence represents cDNA encoding the rabbit MGF.
XX
SO
    Sequence 523 BP; 154 A; 129 C; 142 G; 98 T; 0 other;
 Query Match
                    100.0%; Score 523; DB 24; Length 523;
 Best Local Similarity
                    100.0%; Pred. No. 5.1e-144;
 Matches 523; Conservative
                         0; Mismatches
                                        0; Indels
                                                            0;
                                                      Gaps
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Qy
           1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Db
Qу
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
           61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Db
       121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qy
           Db
       121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
       Qy
           Db
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Qу
           241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qy
           301 AGAAGGAAAGGAAGTACATTTGAAGAACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qy
           361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
           421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Db
       481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
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481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523

Db

```
RESULT 3
AAT84893
ID
     AAT84893 standard; cDNA; 553 BP.
XX
AC
     AAT84893;
XX
DT
     14-APR-1998 (first entry)
XX
DΕ
     Rabbit insulin like growth factor 1 encoding cDNA.
XX
KW
     Insulin like growth factor 1; IGF-1; Ec peptide; muscle disorder;
KW
     heart; neuromuscular disease; primer; ss.
XX
OS
     Oryctolagus cuniculus.
XX
                     Location/Qualifiers
FH
     Kev
\mathbf{FT}
     CDS
                     1..366
FT
                     /*tag= a
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XX
PN
     WO9733997-A1.
XX
PD
     18-SEP-1997.
XX
PF
     11-MAR-1997;
                    97WO-GB00658.
XX
PR
     11-MAR-1996;
                    96GB-0005124.
XX
     (UNLO ) ROYAL FREE HOSPITAL SCHOOL MED.
PA
XX
PΙ
     Goldspink G;
XX
DR
     WPI; 1997-470877/43.
DR
     P-PSDB; AAW23301.
XX
PΤ
     Use of insulin like growth factor I characterised by presence of Ec
РΤ
     peptide - to treat humans or animals, particularly muscle disorders,
PT
     heart conditions or neuromuscular diseases
XX
PS
     Disclosure; Fig 3; 33pp; English.
XX
     A use of insulin like growth factor I (IGF-1) has been developed, and
CC
CC
     is characterised by the presence of the Ec peptide, or a functional
CC
     equivalent, in the treatment or therapy of a human or animal. The IGF-1
     polypeptide can be used to treat muscular disorders, e.g. Duchenne or
CC
CC
     Becker muscular dystrophy, autosomal dystrophies and related progressive
CC
     skeletal muscle weakness and wasting, muscle atrophy in ageing humans,
CC
     spinal cord injury induced muscle atrophy and neuromuscular diseases,
CC
     and cardiac disorders, e.g. diseases where promotion of cardiac muscle
CC
     protein synthesis is a beneficial treatment, cardiomyopathies and acute
CC
     heart failure or insult, specifically myocarditis or myocardial
CC
     infarction. It can also be used to promote bone fracture healing and
CC
     maintenance of bone in old age. The present sequence encodes rabbit
CC
     IGF-1 used in the present specification.
XX
SQ
     Sequence 553 BP; 159 A; 142 C; 147 G; 105 T; 0 other;
```

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Query Match
                   100.0%; Score 523; DB 18;
                                        Length 553;
 Best Local Similarity
                   100.0%; Pred. No. 5.3e-144;
 Matches 523; Conservative
                       0; Mismatches
                                     0; Indels
                                               0;
                                                  Gaps
                                                        0;
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Qу
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Db
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Qy
          91 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 150
Db
       121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
          151 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 210
Db
       Qу
          Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAATGAAGTCTCAGAGG 300
Qy
          Db
       271 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 330
Qу
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
          Dh
       331 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 390
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
          391 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 450
Db
Qy
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
          Db
       451 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 510
       481 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
          511 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 553
Db
RESULT 4
   AAD06398 standard; cDNA; 517 BP.
XX
AC
   AAD06398;
XX
   10-AUG-2001 (first entry)
DT
XX
DE
   Human IGF-I isoform mechano-growth factor (MGF) cDNA.
XX
   Human; IGF-I isoform; Insulin-like Growth Factor-I; MGF;
KW
ΚW
   mechano-growth factor; neurological disorder; neurodegenerative disorder;
KW
   amyotrophic lateral sclerosis; spinal muscular atrophy; muscular atrophy;
KW
   poliomyelitis; post-polio syndrome; toxin; motoneurone disorder;
KW
   nerve damage; autosomal muscular dystrophy; diabetic neuropathy;
```

```
ΚW
     sex-linked muscular dystrophy; peripheral neuropathy;
KW
     Alzheimer's disease; Parkinson's disease; ss.
XX
OS
     Homo sapiens.
XX
FH
     Key
                     Location/Qualifiers
FT
     CDS
                     1..333
FT
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                     /product= "Mechano-growth factor (MGF)"
FT
                     /note= "This region comprises exons 3-6. The CDS does
FT
FT
                     not include start codon"
FT
                     /partial
XX
     WO200136483-A1.
ΡN
XX
PD
     25-MAY-2001.
XX
     15-NOV-2000; 2000WO-GB04354.
PF
XX
PR
                    99GB-0026968.
     15-NOV-1999;
XX
     (UNLO ) UNIV COLLEGE LONDON.
PA
XX
PΙ
     Goldspink G, Johnson I;
XX
DR
     WPI; 2001-355620/37.
DR
     P-PSDB; AAE02447.
XX
PT
     Use of mechano-growth factor, an isoform of Insulin-like Growth
PT
     Factor-I, capable of reducing motoneurone loss, in the manufacture of a
PΤ
     medicament for the treatment of neurological disorder -
XX
PS
     Claim 4; Page 49-50; 66pp; English.
XX
CC
     The present invention relates to use of mechano-growth factor (MGF),
CC
     an Insulin-like Growth Factor-I (IGF-I) isoform in the manufacture of a
CC
    medicament for the treatment of neurological disorder. The MGF is capable
CC
     of reducing motoneurone loss by 20% or greater in response to nerve
CC
     avulsion, and effects motoneurone rescue, preferably adult motoneurone
CC
     rescue. The MGF polynucleotide and polypeptide are useful in the
CC
     manufacture of a medicament for the treatment of a neurological disorder,
CC
     including a disorder of motoneurones and/or neurodegenerative disorder,
CC
     e.g., amyotrophic lateral sclerosis, spinal muscular atrophy, progressive
     spinal muscular atrophy, infantile or juvenile muscular atrophy,
CC
     poliomyelitis or post-polio syndrome, a disorder caused by exposure to a
CC
CC
     toxin, motoneurone trauma, a motoneurone lesion or nerve damage, an
CC
     injury that affects motoneurones, motoneurone loss associated with aging,
CC
     autosomal or sex-linked muscular dystrophy, diabetic neuropathy,
CC
     peripheral neuropathies, Alzheimer's disease and Parkinson's disease.
CC
     The present sequence is human IGF-I isoform MGF cDNA. MGF is a muscle
CC
     isoform having extracellular (Ec) domain, hence also referred as
CC
     IGF-I-Ec. The MGF protein comprises amino acid sequences encoded by
CC
     nucleic acid sequence of IGF-I exons 4, 5 and 6 in the reading frame
CC
     of MGF.
XX
SQ
     Sequence 517 BP; 150 A; 130 C; 139 G; 98 T; 0 other;
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                   89.4%; Score 467.4; DB 22; Length 517;
 Best Local Similarity
                   96.2%;
                         Pred. No. 1.3e-127;
 Matches 501; Conservative
                         0: Mismatches
                                      16: Indels
                                                    Gaps
                                                          2:
Qу
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          Db
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          Db
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          Db
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Qу
          Db
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Qy
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
          Db
       298 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGGAGTGCAGGAAACAAGAACTA 357
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
          Db
       358 CAGGATGTA-GAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 416
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
          417 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 476
Db
       481 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACAT 521
Qу
          477 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACAT 517
Db
RESULT 5
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   AAS16877 standard; cDNA; 517 BP.
XX
   AAS16877;
AC
XX
DT
   25-FEB-2002 (first entry)
XX
DE
   Human mechano-growth factor (MGF) cDNA.
XX
   Human; mechano-growth factor; insulin-like growth factor I; IGF-I; MGF;
KW
KW
   neuroprotective; nerve damage; peripheral nervous system; nerve severing;
   muscle; neurological disorder; motoneuron loss; motorneuron disorder; ss;
KW
KW
   nerve avulsion.
XX
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OS

Homo sapiens.

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XX
FΗ
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                     Location/Qualifiers
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                     /number= 6
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     15-NOV-2001.
XX
PF
     10-MAY-2001; 2001WO-GB02054.
XX
PR
     10-MAY-2000; 2000GB-0011278.
XX
PΑ
     (UNLO ) UNIV COLLEGE LONDON.
     (EGRI-) EAST GRINSTEAD MEDICAL RES TRUST.
PA
XX
PI
     Goldspink G, Terenghi G;
XX
DR
     WPI; 2002-055585/07.
DR
     P-PSDB; AAU10559.
XX
PT
     Use of insulin-like growth factor I (IGF-I) isoform known as
PT
     mechano-growth factor which is encoded by IGF-I exons 4,5,6 and has
PT
     ability to reduce motoneuron loss in response to nerve avulsion, to
PT
     treat nerve damage -
XX
PS
     Claim 11; Fig 5; 65pp; English.
XX
CC
     The invention relates to the use of an insulin-like growth factor I
CC
     (IGF-I) isoform, known as mechano-growth factor (MGF), in the manufacture
CC
     of a medicament for treating nerve damage in the peripheral nervous
CC
     system, or for treating nerve damage by localising MGF at the site of
CC
     damage. The nerve damage may include severing of a nerve. The treatment
CC
     may be combined with another treatment (such as a polypeptide growth
CC
     factor other than MGF) that prevents or diminishes degeneration of the
CC
     target organ (for example, muscle) which the damaged nerve innervates,
CC
     whereby the treatment of the muscle with MGF or a polynucleotide encoding
     MGF prevents or diminishes degeneration. The method is useful for
CC
CC
     treating neurological disorders, preferably motorneuron disorders. These
CC
     methods can reduce motoneuron loss by 20% or greater in response to nerve
CC
     avulsion. This sequence represents cDNA encoding the human MGF.
```

XX

```
Query Match
                   89.4%; Score 467.4; DB 24; Length 517;
 Best Local Similarity
                  96.2%; Pred. No. 1.3e-127;
 Matches 501; Conservative
                         0; Mismatches
                                      16:
                                                          2;
                                         Indels
                                                    Gaps
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Qу
          1 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Db
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Qу
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Db
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Db
       Qу
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Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
          Db
       241 ATGCCCAAGACCCAGAAGTATCAGCCCCCATCTACCAACAAGAACACGAAGTCTCA---G 297
Qу
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
          298 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 357
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qy
          358 CAGGATGTA-GAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 416
Dh
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qy
          417 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 476
Db
       481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACAT 521
Qy
          477 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACAT 517
Db
RESULT 6
AAD06405
   AAD06405 standard; cDNA; 471 BP.
ID
XX
   AAD06405;
AC
XX
DT
   10-AUG-2001 (first entry)
XX
DE
   Rabbit liver-type IGF-I isoform (L.IGF-I) cDNA.
XX
   Rabbit; IGF-I isoform; Insulin-like Growth Factor-I; MGF;
KW
KW
   mechano-growth factor; neurological disorder; neurodegenerative disorder;
KW
   amyotrophic lateral sclerosis; spinal muscular atrophy; muscular atrophy;
   poliomyelitis; post-polio syndrome; toxin; motoneurone disorder;
ΚW
```

```
KW
     nerve damage; autosomal muscular dystrophy; diabetic neuropathy;
     sex-linked muscular dystrophy; peripheral neuropathy;
KW
KW
     Alzheimer's disease; Parkinson's disease; liver; L.IGF-I; ss.
XX
OS
     Oryctolagus cuniculus.
XX
FΗ
     Key
                     Location/Qualifiers
FT
     CDS
                     1..318
FT
                     /*tag= a
FT
                     /product= "Liver-type IGF-I isoform (L.IGF-I)"
FT
                     /transl except= (pos:7..9, aa:Gln)
FT
                     /transl except= (pos:25..27, aa:Gln)
                     /note= "These translation exceptions occur while decoding
FT
FT
                     the alternative version of the protein (AAE02456).
FT
                     The CDS comprises exons 3, 4 and 6 and
                     does not include start codon"
FT
                     /partial
FT
XX
PN
     WO200136483-A1.
XX
     25-MAY-2001.
PD
XX
PF
     15-NOV-2000; 2000WO-GB04354.
XX
PR
     15-NOV-1999;
                    99GB-0026968.
XX
PA
     (UNLO ) UNIV COLLEGE LONDON.
XX
ΡI
     Goldspink G, Johnson I;
XX
DR
     WPI; 2001-355620/37.
DR
     P-PSDB; AAE02452, AAE02456.
XX
PT
     Use of mechano-growth factor, an isoform of Insulin-like Growth
PT
     Factor-I, capable of reducing motoneurone loss, in the manufacture of a
PT
     medicament for the treatment of neurological disorder -
XX
PS
     Disclosure; Page 59-60; 66pp; English.
XX
CC
     The present invention relates to use of mechano-growth factor (MGF),
CC
     an Insulin-like Growth Factor-I (IGF-I) isoform in the manufacture of a
CC
     medicament for the treatment of neurological disorder. The MGF is capable
CC
     of reducing motoneurone loss by 20% or greater in response to nerve
CC
     avulsion, and effects motoneurone rescue, preferably adult motoneurone
CC
     rescue. The MGF polynucleotide and polypeptide are useful in the
CC
     manufacture of a medicament for the treatment of a neurological disorder,
CC
     including a disorder of motoneurones and/or neurodegenerative disorder,
CC
     e.g., amyotrophic lateral sclerosis, spinal muscular atrophy, progressive
CC
     spinal muscular atrophy, infantile or juvenile muscular atrophy,
CC
     poliomyelitis or post-polio syndrome, a disorder caused by exposure to a
CC
     toxin, motoneurone trauma, a motoneurone lesion or nerve damage, an
CC
     injury that affects motoneurones, motoneurone loss associated with aging,
CC
     autosomal or sex-linked muscular dystrophy, diabetic neuropathy,
CC
     peripheral neuropathies, Alzheimer's disease and Parkinson's disease.
CC
     The present sequence is rabbit liver-type IGF-I isoform (L.IGF-I) cDNA.
CC
     The L.IGF-I protein comprises amino acid sequences encoded by
CC
     nucleic acid sequence of IGF-I exons 4 and 6.
```

```
XX
SO
```

Sequence 471 BP; 132 A; 118 C; 131 G; 90 T; 0 other;

```
Query Match
                  78.2%; Score 409; DB 22; Length 471;
 Best Local Similarity 90.1%; Pred. No. 2e-110;
 Matches 471; Conservative
                       0; Mismatches
                                  0; Indels
                                                       1;
        1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
          Db
        1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
          61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qy
          121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
       Qу
          Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAATGAAGTCTCAGAGG 300
Qу
          Db
       241 ATGCCCAAGACTCAG----- 255
Qу
       301 AGAAGGAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
               256 -----AAGGAAGTACATTTGAAGAACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 308
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qy
          309 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 368
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
          369 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 428
Db
       481 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
          429 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 471
Dh
RESULT 7
   AAS16884 standard; cDNA; 471 BP.
XX
AC
   AAS16884;
XX
DΤ
   25-FEB-2002 (first entry)
XX
DE
   Rabbit insulin-like growth factor I liver-type isoform (L.IGF-I) cDNA.
XX
KW
   Rabbit; mechano-growth factor; insulin-like growth factor I; IGF-I; MGF;
KW
   neuroprotective; nerve damage; peripheral nervous system; nerve severing;
KW
   muscle; neurological disorder; motoneuron loss; motorneuron disorder; ss;
```

```
KW
     nerve avulsion; insulin-like growth factor I liver-type isoform; L.IGF-I;
XX
OS
     Oryctolagus cuniculus.
XX
FΗ
                     Location/Qualifiers
     Key
FT
     CDS
                     1..318
FT
                     /*tag=
FΤ
                     /product= "Rabbit L.IGF-I"
FT
                     /partial
FΤ
                     /note= "No start codon"
FT
                     1..75
     exon
FT
                     /*tag= b
                     /number= exon 3
FT
FT
                     76..258
     exon
FT
                     /*tag= c
                     /number= exon 4
FT
                     259..315
FT
     exon
FT
                     /*tag= d
FT
                     /number= exon 6
XX
PN
     WO200185781-A2.
XX
ΡĎ
     15-NOV-2001.
XX
ΡF
     10-MAY-2001; 2001WO-GB02054.
XX
PR
     10-MAY-2000; 2000GB-0011278.
XX
PA
     (UNLO ) UNIV COLLEGE LONDON.
     (EGRI-) EAST GRINSTEAD MEDICAL RES TRUST.
PΑ
XX
PΙ
     Goldspink G, Terenghi G;
XX
     WPI; 2002-055585/07.
DR
DR
     P-PSDB; AAU10564.
XX
PT
     Use of insulin-like growth factor I (IGF-I) isoform known as
PT
     mechano-growth factor which is encoded by IGF-I exons 4,5,6 and has
PT
     ability to reduce motoneuron loss in response to nerve avulsion, to
РΤ
     treat nerve damage -
XX
PS
     Disclosure; Fig 10; 65pp; English.
XX
CC
     The invention relates to the use of an insulin-like growth factor I
CC
     (IGF-I) isoform, known as mechano-growth factor (MGF), in the manufacture
CC
     of a medicament for treating nerve damage in the peripheral nervous
CC
     system, or for treating nerve damage by localising MGF at the site of
CC
     damage. The nerve damage may include severing of a nerve. The treatment
CC
    may be combined with another treatment (such as a polypeptide growth
CC
     factor other than MGF) that prevents or diminishes degeneration of the
CC
     target organ (for example, muscle) which the damaged nerve innervates,
CC
     whereby the treatment of the muscle with MGF or a polynucleotide encoding
CC
    MGF prevents or diminishes degeneration. The method is useful for
CC
     treating neurological disorders, preferably motorneuron disorders. These
CC
     methods can reduce motoneuron loss by 20% or greater in response to nerve
CC
     avulsion. This sequence represents cDNA encoding the rabbit insulin-like
CC
     growth factor I liver-type isoform (L.IGF-I) used in experiments on
```

```
CC
   motoneuron loss.
XX
SQ
   Sequence 471 BP; 132 A; 118 C; 131 G; 90 T; 0 other;
 Query Match
                   78.2%; Score 409; DB 24; Length 471;
 Best Local Similarity
                  90.1%; Pred. No. 2e-110;
 Matches 471; Conservative
                        0; Mismatches
                                     0;
                                       Indels
                                              52:
                                                  Gaps
                                                        1:
        1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
          Db
        1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qy
          Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
          Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
       Qy
          Db
       Qy
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
          111111111111
Db
       241 ATGCCCAAGACTCAG----- 255
Qу
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
               Db
       256 -----AAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 308
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
          309 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 368
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
          369 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 428
Db
       481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
          429 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 471
Db
RESULT 8
AAD06399
ID
   AAD06399 standard; cDNA; 539 BP.
XX
AC
   AAD06399;
XX
DΤ
   10-AUG-2001 (first entry)
XX
DE
   Rat IGF-I isoform mechano-growth factor (MGF) cDNA.
XX
KW
   Rat; IGF-I isoform; Insulin-like Growth Factor-I; MGF;
KW
   mechano-growth factor; neurological disorder; neurodegenerative disorder;
```

```
KW
     amyotrophic lateral sclerosis; spinal muscular atrophy; muscular atrophy;
KW
     poliomyelitis; post-polio syndrome; toxin; motoneurone disorder;
KW
     nerve damage; autosomal muscular dystrophy; diabetic neuropathy;
KW
     sex-linked muscular dystrophy; peripheral neuropathy;
     Alzheimer's disease; Parkinson's disease; ss.
KW
XX
OS
     Rattus sp.
XX
FΗ
     Key
                     Location/Qualifiers
     CDS
FT
                     1..336
FT
                     /*tag= a
FT
                     /product= "Mechano-growth factor (MGF)"
                     /note= "This region comprises exons 3-6. The CDS does
FT
FT
                     not include start codon"
FT
                     /partial
XX
PN
     WO200136483-A1.
XX
PD
     25-MAY-2001.
XX
     15-NOV-2000; 2000WO-GB04354.
PF
XX
PR
     15-NOV-1999;
                    99GB-0026968.
XX
PA
     (UNLO ) UNIV COLLEGE LONDON.
XX
PI
     Goldspink G, Johnson I;
XX
     WPI: 2001-355620/37.
DR
DR
     P-PSDB; AAE02448.
XX
     Use of mechano-growth factor, an isoform of Insulin-like Growth
PT
     Factor-I, capable of reducing motoneurone loss, in the manufacture of a
PT
PT
     medicament for the treatment of neurological disorder -
XX
PS
     Claim 4; Page 51-52; 66pp; English.
XX
CC
     The present invention relates to use of mechano-growth factor (MGF),
CC
     an Insulin-like Growth Factor-I (IGF-I) isoform in the manufacture of a
CC
     medicament for the treatment of neurological disorder. The MGF is capable
CC
     of reducing motoneurone loss by 20% or greater in response to nerve
CC
     avulsion, and effects motoneurone rescue, preferably adult motoneurone
CC
     rescue. The MGF polynucleotide and polypeptide are useful in the
CC
     manufacture of a medicament for the treatment of a neurological disorder,
     including a disorder of motoneurones and/or neurodegenerative disorder,
CC
CC
     e.g., amyotrophic lateral sclerosis, spinal muscular atrophy, progressive
CC
     spinal muscular atrophy, infantile or juvenile muscular atrophy,
CC
     poliomyelitis or post-polio syndrome, a disorder caused by exposure to a
     toxin, motoneurone trauma, a motoneurone lesion or nerve damage, an
CC
CC
     injury that affects motoneurones, motoneurone loss associated with aging,
CC
     autosomal or sex-linked muscular dystrophy, diabetic neuropathy,
CC
     peripheral neuropathies, Alzheimer's disease and Parkinson's disease.
     The present sequence is rat IGF-I isoform MGF cDNA. MGF is a muscle
CC
CC
     isoform having extracellular (Ec) domain, hence also referred as
CC
     IGF-I-Ec. The MGF protein comprises amino acid sequences encoded by
CC
     nucleic acid sequence of IGF-I exons 4, 5 and 6 in the reading frame
CC
     of MGF.
```

```
XX
SO
```

Sequence 539 BP; 161 A; 136 C; 139 G; 103 T; 0 other;

```
Query Match
                    68.2%; Score 356.8; DB 22; Length 539;
 Best Local Similarity
                    82.3%; Pred. No. 5.1e-95;
 Matches 436; Conservative
                         0; Mismatches 87; Indels
                                                           2;
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
           Db
         1 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGACGCTCTTCAGTTCGTGTGTGGACCA 60
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qy
           61 AGGGGCTTTTACTTCAACAAGCCCACAGTCTATGGCTCCAGCATTCGGAGGGCACCACAG 120
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           Db
       121 ACGGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
       Qу
           181 TGTGTCCGCTGCAAGCCTACAAAGTCAGCTCGTTCCATCCGGGCCCAGCGCCACACTGAC 240
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
           Db
       241 ATGCCCAAGACTCAGAAGTCCCAGCCCCTATCGACACACAAGAAAAGGAAGCTGCAAAGG 300
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
           301 AGAAGGAAAGGAAGTACACTTGAAGAACAAGTAGAGGAAGTGCAGGAAACAAGACCTA 360
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qy
          361 CAGAATGTAGGAGGAGCCTCCCGAGGAACAGAAAATGCCACGTCACCGCAAGATCCTTTG 420
Db
Qу
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA----AAAAATAAGTTTGATC 474
          \Pi
                    1
                        11111 1111 111
                                            421 CTGCTTGAGCAACCTGCAAAACATCGGAACACCTGCCAAATATCAATAATGAGTTCAATA 480
Db
       475 ACATTTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           481 TCATTTCAGAGATGGGCATTTCCCTCAATGAAATACACAAGTAAACATTC 530
Db
RESULT 9
AAS16878
   AAS16878 standard; cDNA; 539 BP.
XX
AC
   AAS16878;
XX
DT
   25-FEB-2002 (first entry)
XX
   Rat mechano-growth factor (MGF) cDNA.
DE
XX
KW
    Rat; mechano-growth factor; insulin-like growth factor I; IGF-I; MGF;
KW
    neuroprotective; nerve damage; peripheral nervous system; nerve severing;
KW
   muscle; neurological disorder; motoneuron loss; motorneuron disorder; ss;
```

```
KW
     nerve avulsion.
XX
OS
     Rattus sp.
XX
FH
                     Location/Qualifiers
     Key
FT
     CDS
                     1..336
FT
                     /*tag= a
FT
                     /product= "Rat MGF"
FT
                     /partial
FT
                     /note= "No start codon"
FT
     exon
                     1..75
FT
                     /*tag= b
                     /number= exon 3
FT
FT
                     76..258
     exon
                     /*tag= c
FT
FT
                     /number= exon 4
                     259..309
FT
     exon
FT
                     /*tag= d
FT
                     /number= exon 5
FT
                     310..333
     exon
                     /*tag= e
FT
                     /number= exon 6
FT
XX
    WO200185781-A2.
ΡN
XX
PD
     15-NOV-2001.
XX
    10-MAY-2001; 2001WO-GB02054.
PF
XX
PR
     10-MAY-2000; 2000GB-0011278.
XX
     (UNLO ) UNIV COLLEGE LONDON.
PA
PA
     (EGRI-) EAST GRINSTEAD MEDICAL RES TRUST.
XX
PI
     Goldspink G, Terenghi G;
XX
     WPI; 2002-055585/07.
DR
     P-PSDB; AAU10560.
DR
XX
PT
     Use of insulin-like growth factor I (IGF-I) isoform known as
     mechano-growth factor which is encoded by IGF-I exons 4,5,6 and has
PT
PΤ
     ability to reduce motoneuron loss in response to nerve avulsion, to
     treat nerve damage
PT
XX
PS
     Disclosure; Fig 6; 65pp; English.
XX
CC
     The invention relates to the use of an insulin-like growth factor I
CC
     (IGF-I) isoform, known as mechano-growth factor (MGF), in the manufacture
CC
     of a medicament for treating nerve damage in the peripheral nervous
     system, or for treating nerve damage by localising MGF at the site of
CC
CC
     damage. The nerve damage may include severing of a nerve. The treatment
CC
     may be combined with another treatment (such as a polypeptide growth
CC
     factor other than MGF) that prevents or diminishes degeneration of the
     target organ (for example, muscle) which the damaged nerve innervates,
CC
CC
     whereby the treatment of the muscle with MGF or a polynucleotide encoding
CC
     MGF prevents or diminishes degeneration. The method is useful for
     treating neurological disorders, preferably motorneuron disorders. These
CC
```

```
CC
    methods can reduce motoneuron loss by 20% or greater in response to nerve
CC
    avulsion. This sequence represents cDNA encoding the rat MGF.
XX
SO
    Sequence 539 BP; 161 A; 136 C; 139 G; 103 T; 0 other;
                    68.2%; Score 356.8; DB 24; Length 539;
 Best Local Similarity
                    82.3%; Pred. No. 5.1e-95;
 Matches 436; Conservative
                         0; Mismatches 87; Indels
                                                 7; Gaps
                                                           2:
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Qy
           Db
         1 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGACGCTCTTCAGTTCGTGTGTGGACCA 60
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qy
          Db
        61 AGGGGCTTTTACTTCAACAAGCCCACAGTCTATGGCTCCAGCATTCGGAGGGCACCACAG 120
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qy
           Db
       121 ACGGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
       Qy
          Db
       181 TGTGTCCGCTGCAAGCCTACAAAGTCAGCTCGTTCCATCCGGGCCCAGCGCCACACTGAC 240
Qу
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAATGAAGTCTCAGAGG 300
          241 ATGCCCAAGACTCAGAAGTCCCAGCCCCTATCGACACACAAGAAAAGGAAGCTGCAAAGG 300
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
          301 AGAAGGAAAGGAAGTACACTTGAAGAACAAGTAGAGGAAGTGCAGGAAACAAGACCTA 360
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qy
          Db
       361 CAGAATGTAGGAGGAGCCTCCCGAGGAACAGAAAATGCCACGTCACCGCAAGATCCTTTG 420
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA-----AAAAATAAGTTTGATC 474
Qу
          \Pi
                    1
                        Db
       421 CTGCTTGAGCAACCTGCAAAACATCGGAACACCTGCCAAATATCAATAATGAGTTCAATA 480
       475 ACATTTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           Db
       481 TCATTTCAGAGATGGGCATTTCCCTCAATGAAATACACAAGTAAACATTC 530
RESULT 10
ABV76185
ID
   ABV76185 standard; cDNA; 651 BP.
XX
AC
   ABV76185;
XX
   07-MAR-2003 (first entry)
DT
XX
DE
   Mouse insulin-like growth factor IB cDNA.
XX
KW
   Insulin-like growth factor IB; IGF-IB; mouse; mRNA; assay;
```

```
KW
     nucleic acid detection; gene; ss.
XX
OS
     Mus musculus.
XX
FH
     Key
                     Location/Qualifiers
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     CDS
                     73..474
FT
                     /*tag= a
FT
                     /product= "IGF-IB"
XX
PN
     WO200297390-A2.
XX
     05-DEC-2002.
PD
XX
PF
     31-MAY-2002; 2002WO-SE01056.
XX
     01-JUN-2001; 2001SE-0001934.
PR
XX
PΑ
     (BIOV-) BIOVITRUM AB.
XX
PI
     Parrow V, Rosengren L;
XX
DR
     WPI; 2003-129529/12.
XX
PT
     Quantitating a target nucleic acid in a sample comprises immobilizing,
РΤ
     on a solid support, a sample comprising a target nucleic acid, and
PT
     detecting and quantitating signals generated from the antisense and
PT
     sense probes -
XX
PS
     Example 1; Page 16-17; 18pp; English.
XX
CC
     The present sequence is that of cDNA encoding murine insulin-like
     growth factor 1B (IGF-IB). The cDNA was used in an example of the
CC
CC
     method of the invention to generate probes for determination of
CC
     IGF-IB RNA. The method comprises a quantitative hybridisation
CC
     assay for analysis of mRNA in a target nucleic acid (TNA) sample.
CC
     It involves: (i) immobilising the TNA sample on a solid support;
CC
     (ii) contacting a labelled antisense probe to a first portion of the
CC
     TNA, and a labelled sense probe to a second portion of the TNA;
CC
     (iii) detecting and quantitating the signals generated from the
CC
     hybridised probes; and (iv) determining the value represented by
CC
     the antisense probe signal minus the sense probe signal, the value
CC
     being proportional to the amount of mRNA in the TNA sample. In an
CC
     example of the method, a cDNA clone containing 60 nucleotides from
CC
     exon 2 and 179 nucleotides from exon 3 of the mouse IGF-IB gene was
CC
     cloned into pGEN-4Z vector. Linearisation of the plasmid with
CC
     EcoRI allowed transcription of a 250-nucleotide antisense probe
CC
     using T7 polymerase. Linearisation with HindIII allowed
CC
     transcription of a sense probe of similar length using SP6
CC
     polymerase (see ABV76186). The probes were purified and used to
CC
     determine IGF-I RNA in mouse hepatocytes and also in rat hepatocytes.
XX
SO
     Sequence 651 BP; 193 A; 185 C; 149 G; 124 T; 0 other;
  Query Match
                          66.8%; Score 349.4; DB 25; Length 651;
```

82.8%; Pred. No. 8.3e-93; 0; Mismatches 81; Indels 7; Gaps 2;

Matches 425; Conservative

Best Local Similarity

```
Qу
        1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
          Db
       139 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGACCG 198
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
          199 AGGGGCTTTTACTTCAACAAGCCCACAGGCTATGGCTCCAGCATTCGGAGGGCACCTCAG 258
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
          259 ACAGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGACTGGAGATGTAC 318
Db
       Qу
          319 TGTGCCCCACTGAAGCCTACAAAAGCAGCCCGCTCTATCCGTGCCCAGCGCCACACTGAC 378
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
          Db
Qy
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
          439 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGAAGTGCAGGAAACAAGACCTA 498
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
          Db
       499 CAGAATGTAGGAGGAGCCTCCCACGGAGCAGAAAATGCCACATCACCGCAGGATCCTTTG 558
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA----AAAAATAAGTTTGATC 474
Qу
          11
                       11 111111
       559 CTGCTTGAGCAACCTGCAAAACATCGAAACACCTACCAAATAACAATAATAAGTCCAATA 618
Db
Qу
       475 ACATTTCAAAGAT-GGCATTTCCCCCAATGAAA 506
          619 ACATTACAAAGATGGGCATTTCCCCCAATGAAA 651
Db
RESULT 11
AAN70436
   AAN70436 standard; cDNA; 818 BP.
ID
XX
AC
   AAN70436;
XX
DT
   25-MAR-2003
             (updated)
DT
             (first entry)
   05-APR-1991
XX
DE
   Sequence encoding insulin-like growth factor 1A (IGF-1A).
XX
KW
   Growth promoter; lactation enhancer; cell proliferation; ss.
XX
OS
   Homo sapiens.
XX
PN
   EP229750-A.
XX
PD
   22-JUL-1987.
XX
PF
              87EP-0870001.
   06-JAN-1987;
```

```
XX
PR
    20-NOV-1986;
                86US-0929671.
PR
    07-JAN-1986;
                 86US-0816662.
XX
PA
    (UNIW ) UNIV WASHINGTON.
XX
PΙ
    Krivi GG, Rotwein PS;
XX
DR
    WPI; 1987-200203/29.
XX
PT
    New pre-pro-insulin-like growth factor-1 protein - obtd. by
PΤ
    recombinant DNA procedures for use as growth promoters for
PT
    enhancing lactation, for stimulating cell proliferation etc.
XX
PS
    Example; Fig 5; 59pp; English.
XX
CC
    A 42 base oligonucleotide corresponding to the DNA sequence encoding
CC
    amino acids 10 to 23 of mature human IGF-I was synthesized (AAN70437).
CC
    The radiolabeled 42 mer was then employed to screen for IGF-I
CC
    containing DNA sequences in a human liver cDNA library. Insulin-
CC
    like growth factors-1A and -1B cDNAs were isolated from a human cDNA
CC
    library by using lambdagt 11 (AAN70435, AAN70436). The human IGF-1
CC
    genomic gene was isolated and mapped. It encodes at least two
CC
    preproinsulin-like growth factor-1 proteins. An essentially pure
CC
    proproinsulin-like growth factor-1 protein comprising the sequence
CC
    of amino acids shown in Figure six is claimed (AAP70277).
CC
    (Updated on 25-MAR-2003 to correct PA field.)
XX
    Sequence 818 BP; 232 A; 186 C; 187 G; 213 T; 0 other;
SQ
 Query Match
                     63.9%; Score 334.4; DB 8; Length 818;
 Best Local Similarity
                     84.6%; Pred. No. 2.4e-88;
 Matches 445; Conservative
                           0; Mismatches
                                         26; Indels
                                                     55;
                                                         Gaps
                                                                4;
Qy
          1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
           Db
        203 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 262
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCCACCTCAG 120
Qy
           Db
        263 AGGGGCTTTTATTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 322
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           Db
        323 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 382
        Qy
           383 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 442
Db
        241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qy
           Db
        443 ATGCCCAAGACCCAG----- 457
Qу
        301 AGAAGGAAAGGAAGTACATTTGAAGAACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
                 458 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 510
Db
```

```
361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
             1111 11111111111
Db
         511 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 570
Qу
         421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
             571 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAATAAGTTTGATAACAT 630
Db
         479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
             631 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 676
Db
RESULT 12
ABT11091
ID
    ABT11091 standard; cDNA; 7260 BP.
XX
AC
    ABT11091;
XX
DT
    04-DEC-2002 (first entry)
XX
DE
    Human breast cancer associated coding sequence SEQ ID NO: 1225.
XX
ΚW
    Human; breast specific gene; breast cancer; differential expression;
    cytostatic; gene therapy; gene; ss.
KW
XX
OS
    Homo sapiens.
XX
PN
    WO200259271-A2.
XX
    01-AUG-2002.
PD
XX
PF
    25-JAN-2002; 2002WO-US02176.
XX
PR
    25-JAN-2001; 2001US-263757P.
    25-APR-2001; 2001US-286090P.
PR
PR
    23-MAY-2001; 2001US-292517P.
XX
    (GENE-) GENE LOGIC INC.
PA
XX
PΙ
    Orr MS, Nation M, Diggans JC,
                                   Zeng W;
XX
DR
    WPI; 2002-674803/72.
XX
PT
    Diagnosing breast cancer in a patient comprises detecting the level of
PT
    gene expression in cell or tissue samples, where a differential gene
PT
    expression is indicative of breast cancer -
XX
PS
    Claim 1; SEQ ID NO 1225; 260pp + Sequence Listing; English.
XX
CC
    The present invention relates to methods of diagnosing breast cancer in a
CC
    patient, which comprise detecting the level of expression in a tissue
CC
    sample of two or more genes selected from those shown in ABT09867-
CC
    ABT11112, where a differential expression of the genes indicates breast
CC
    cancer. The methods are useful in diagnosing, treating, detecting the
    progression, and in monitoring treatment of breast cancer in patients.
CC
```

```
CC
   The methods are also useful as a screening tool for agents that modulate
CC
    the onset or progression of breast cancer. The breast cancer genes may be
CC
    used as diagnostic markers for the prediction or identification of the
CC
    malignant state of breast tissue, for confirming the type and progression
    of cancer, and for drug screening and assays. The present sequence is a
CC
CC
    coding sequence of the invention.
CC
   Note: The sequence data for this patent did not form part of the printed
CC
    specification, but was obtained in electronic format directly from WIPO
   at ftp.wipo.int/pub.published pct sequences.
CC
XX
SQ
    Sequence 7260 BP; 2330 A; 1415 C; 1240 G; 2275 T; 0 other;
                    63.9%; Score 334.4; DB 24; Length 7260;
 Query Match
 Best Local Similarity
                    84.6%; Pred. No. 5.7e-88;
 Matches 445; Conservative
                         0; Mismatches
                                      26; Indels
                                                  55; Gaps
                                                            4;
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qy
           Db
       311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qy
           Db
       371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
QУ
           Db
       431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Qу
       491 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 550
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
           551 ATGCCCAAGACCCAG----- 565
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
                566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
       421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
           679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAATAAGTTTGATAACAT 738
Db
       479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qy
           Db
       739 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 784
```

## RESULT 13 ABK84583

ID ABK84583 standard; cDNA; 7260 BP.

```
AC
     ABK84583;
XX
DT
     14-AUG-2002 (first entry)
XX
DΕ
     Human cDNA differentially expressed in granulocytic cells #1154.
XX
KW
     Human; ss; granulocytic cell; DNA chip; bacterial infection;
KW
     viral infection; parasitic infection; protozoal infection;
KW
     fungal infection; sterile inflammatory disease; psoriasis;
KW
     rheumatoid arthritis; qlomerulonephritis; asthma; thrombosis;
KW
     cardiac reperfusion injury; renal reperfusion injury; ARDS;
KW
     adult respiratory distress syndrome; inflammatory bowel disease;
KW
     Crohn's disease; ulcerative colitis; periodontal disease;
KW
     granulocyte activation; chronic inflammation; allergy.
XX
OS
     Homo sapiens.
XX
     WO200228999-A2.
PN
XX
     11-APR-2002.
PD
XX
PF
     03-OCT-2001; 2001WO-US30821.
XX
PR
     03-OCT-2000; 2000US-237189P.
XX
PA
     (GENE-) GENE LOGIC INC.
XX
PΙ
     Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;
XX
DR
    WPI; 2002-435328/46.
XX
PT
     Detecting granulocyte activation by detecting differential expression
PΤ
     of genes associated with granulocyte activation, which serves as
PT
     diagnostic markers that is useful for monitoring disease states and
PT
    drug toxicity
XX
PS
     Claim 1; SEQ ID No 1154; 114pp; English.
XX
CC
     The invention relates to detecting (M1) granulocyte (GC) activation
CC
     (GCA), by detecting the level of expression of gene(s) (Gs) identified by
CC
     DNA chip analysis as given in the specification, and comparing
CC
     the expression level to an expression level in an unactivated
CC
     GC, where differential expression of Gs is indicative of GCA.
CC
    Also included are modulating (M2) GA by contacting GC with an agent
CC
     that alters the expression of at least one gene in Gs; (2) screening (M3)
CC
     for an agent capable of modulating GCA or an inflammation (especially
CC
     chronic) in a tissue, an allergic response in a subject, exposure of a
CC
     subject to a pathogen or sterile inflammatory disease using the
CC
     gene expression profile; (3) detecting (M4) an inflammation (especially
CC
     chronic) in a tissue, an allergic response in a subject, exposure of a
CC
     subject to a pathogen or sterile inflammatory disease, by detecting the
CC
     level of expression in a sample of the tissue of gene(s) from Gs, where
CC
     the level of expression of the gene is indicative of inflammation;
CC
     (4) treating (M5) an inflammation (especially chronic) or in a tissue,
CC
     an allergic response in a subject, exposure of a subject to a pathogen
CC
     or sterile inflammatory disease, by contacting a tissue having
CC
     inflammation with an agent that modulates the expression of gene(s)
```

```
CC
    from Gs in the tissue. M1 is useful for detecting GCA; M2 is useful for
CC
    modulating GA; M3 is useful for screening an agent capable of modulating
CC
    GCA preferably in an inflammation in a tissue; M4 is useful for
CC
    detecting an inflammation (especially chronic) in a tissue, an allergic
CC
    response in a subject, exposure of a subject to a pathogen or sterile
CC
    inflammatory disease (e.g. psoriasis, rheumatoid arthritis,
CC
    glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal
CC
    reperfusion injury, ARDS, adult respiratory distress syndrome,
CC
    inflammatory bowel disease, Crohn's disease, ulcerative colitis,
CC
    periodontal disease; also bacterial infection, viral infection,
CC
    parasitic infection, protozoal infection, fungal infection and M5 is
CC
    useful for treating one of the above conditions. The present
CC
    sequence represents a gene differentially expressed in granulocytes.
CC
    Note: The sequence data for this patent did not form part
CC
    of the printed specification, but was obtained in electronic
CC
    format directly from WIPO at
CC
    ftp.wipo.int/pub/published pct sequences.
XX
SO
    Sequence 7260 BP; 2330 A; 1415 C; 1240 G; 2275 T; 0 other;
 Query Match
                     63.9%; Score 334.4; DB 24; Length 7260;
 Best Local Similarity 84.6%; Pred. No. 5.7e-88;
 Matches 445; Conservative 0; Mismatches 26; Indels
                                                    55; Gaps
                                                               4:
Qу
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
           Db
        311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
Qу
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCCACCTCAG 120
           Db
        371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
        Qу
           491 TGCGCACCCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 550
Db
        241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
           11111111111 111
        551 ATGCCCAAGACCCAG----- 565
Db
        301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
                 566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Db
        361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
        421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qy
           Db
        679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
```

479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523

Qу

#### 739 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 784

RESULT 14 ABN97244 ID ABN97244 standard; DNA; 7260 BP. XX AC ABN97244; XX 13-AUG-2002 (first entry) DT XX Gene #3742 used to diagnose liver cancer. DΕ XX Gene; liver cancer; ds; hepatocellular carcinoma; hepatotropic; KW KW metastatic liver tumour; cytostatic; expression profile; disease state; KW disease progression; drug toxicity; drug efficacy; drug metabolism. XX OS Homo sapiens. XX WO200229103-A2. PN XX PD 11-APR-2002. XX PF 02-OCT-2001; 2001WO-US30589. XX 02-OCT-2000; 2000US-237054P. PR XX PA (GENE-) GENE LOGIC INC. XX PΙ Horne D, Alvares C, Peres-Da-Silva S, Vockley JG; XX DR WPI; 2002-426119/45. XX PTDiagnosing and detecting the progression of liver cancer, PThepatocellular carcinoma or metastatic liver tumor in a patient, involves detecting the level of expression of two or more genes in a PТ liver tissue sample XX PS Claim 1; SEQ ID NO 3742; 298pp; English. XX CC The invention relates to a novel method for diagnosing and detecting the CC progression of liver cancer, hepatocellular carcinoma or metastatic liver CC tumour in a patient, and differentiating metastatic liver cancer from CC hepatocellular carcinoma in a patient, involving detecting the level of CC expression of two or more genes represented in ABN93503-ABN97455 in a CC tissue sample. The method of the invention has hepatotropic, and CC cytostatic activity. The method is useful for diagnosing and detecting CC the progression of liver cancer, hepatocellular carcinoma and metastatic CC liver carcinoma in a patient. The method is useful for identifying CC expression profiles which serve as useful diagnostic markers as well as CC markers that can be used to monitor disease states, disease progression, CC drug toxicity, drug efficacy and drug metabolism. CC Note: The sequence data for this patent did not form part of the printed CC specification, but was obtained in electronic format directly from WIPO CCat ftp.wipo.int/pub/published pct sequences. XX

```
Query Match
                    63.9%; Score 334.4; DB 24; Length 7260;
 Best Local Similarity
                    84.6%; Pred. No. 5.7e-88;
 Matches 445; Conservative
                         0; Mismatches
                                      26; Indels
                                                 55; Gaps
                                                           4;
Qy
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
           Db
       311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
Qy
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
           Db
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Qу
       121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
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Db
Ov
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Qy
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           11111111111111
Db
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Qy
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                Db
       566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
       421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qy
           11114411 114141414 1411 14114 11144 111 111414 111
       679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
Db
       479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qy
           739 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 784
Db
RESULT 15
ABK64812
ID
   ABK64812 standard; DNA; 7260 BP.
XX
AC
   ABK64812;
XX
DT
   18-JUN-2002 (first entry)
XX
DE
   Human benign prostatic hyperplasia gene #707.
XX
KW
   Human; benign prostatic hyperplasia; BPH; prostate cancer; gene; ds.
XX
   Homo sapiens.
OS
XX
```

```
PN
    WO200212440-A2.
XX
PD
     14-FEB-2002.
XX
PF
     07-AUG-2001; 2001WO-US24708.
XX
PR
     07-AUG-2000; 2000US-223323P.
PR
     05-JUN-2001; 2001US-0873319.
XX
     (GENE-) GENE LOGIC INC.
PA
     (NISB ) JAPAN TOBACCO INC.
PA
XX
PΙ
    Munger WE, Kulkarni P, Getzenberg RH, Waga I, Yamamoto J;
XX
DR
    WPI; 2002-257476/30.
XX
PΨ
     Identifying drugs for and diagnosing benign prostatic hyperplasia, by
PT
     detecting expression levels of one or more genes in prostate cells from
РΤ
     patient that are differentially regulated compared to normal prostate
РΤ
    cells -
XX
PS
    Disclosure; Page 391-393; 444pp; English.
XX
CC
    The invention relates to a method of diagnosing (I) the onset or
CC
    progression of benign prostatic hyperplasia (BPH), or screening (II) for
CC
    or identifying an agent that modulates the onset or progression of BPH.
CC
    The method is based on changes in gene expression in BPH tissue isolated
CC
     from patients exhibiting different clinical states of prostate
CC
     hyperplasia as compared to normal prostate tissue. (I) comprises
CC
     detecting the expression levels of one or more genes in prostate cells
CC
     from the subject that are differentially regulated compared to normal
CC
    prostate cells. (II) comprises preparing a first gene expression profile
CC
     of BPH cells or BPH-like cell population, exposing the cells to the
CC
     agent, preparing a second gene expression profile of the agent exposed
CC
     cells, and comparing the first and second gene expression profiles.
CC
     (I) is useful for diagnosing the onset or progression of BPH. (II) is
CC
    useful for identifying an agent that modulates the onset or progression
CC
    of BPH. The methods are useful to present information identifying
CC
    the expression level in a tissue or cells, by comparing the expression
CC
     level of genes given in the specification in the tissue or cells to the
CC
    level of expression of gene in the database, and displaying the
CC
     expression levels of at least one gene in the tissue or cell sample
CC
     compared to the expression level in BPH. Agents using (II) are useful for
CC
     treating BPH or prostate cancer. ABK64106-ABK64860 represent human
CC
    benign prostatic hyperplasia gene sequences of the invention.
XX
     Sequence 7260 BP; 2330 A; 1415 C; 1240 G; 2275 T; 0 other;
SQ
                         63.9%; Score 334.4; DB 24; Length 7260;
  Query Match
                         84.6%; Pred. No. 5.7e-88;
  Best Local Similarity
                                0; Mismatches
 Matches 445; Conservative
                                                 26; Indels
                                                                            4;
Qу
            1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
              Db
         311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
          61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qy
```

371	${\tt AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG}$	430
121		180
431		490
181		240
491		550
241		300
551	* * * * * * * * * * * * * * * * * * * *	565
301		360
566		618
361		420
619		678
421		478
679		738
479	<del></del>	
739		
	121 431 181 491 241 551 301 566 361 619 421 679 479	

Search completed: December 13, 2003, 06:03:51 Job time : 211.995 secs

### GenCore version 5.1.6 Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 13, 2003, 06:03:55; Search time 48.3585 Seconds

(without alignments)

4773.589 Million cell updates/sec

Title:

Sequence:

US-09-852-261-5

Perfect score: 523

1 ggaccggagacgctctgcgg.....aaatacacaagtaaacattc 523

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched:

569978 segs, 220691566 residues

Total number of hits satisfying chosen parameters: 1139956

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents NA:\*

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- 2: /cgn2\_6/ptodata/1/ina/5B\_COMB.seq:\*
- 3: /cgn2\_6/ptodata/1/ina/6A\_COMB.seq:\*
- 4: /cgn2 6/ptodata/1/ina/6B COMB.seq:\*
- 5: /cgn2 6/ptodata/1/ina/PCTUS COMB.seq:\*
- 6: /cgn2 6/ptodata/1/ina/backfiles1.seq:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

### SUMMARIES

		*				
Result		Query				
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1	523	100.0	553	3	US-09-142-583A-3	Sequence 3, Appli
2	523	100.0	553	3	US-09-142-583A-5	Sequence 5, Appli
3	332.8	63.6	777	3	US-09-142-583A-10	Sequence 10, Appl
4	331.2	63.3	622	6	5405942-2	Patent No. 5405942
5	274.6	52.5	5707	2	US-08-472-809B-8	Sequence 8, Appli
6	274.6	52.5	6345	2	US-08-472-809B-7	Sequence 7, Appli
7	234.4	44.8	357	6	5405942-13	Patent No. 5405942
8	232.8	44.5	357	6	5405942-9	Patent No. 5405942
9	191.4	36.6	210	6	5405942-7	Patent No. 5405942
10	191.4	36.6	210	6	5405942-11	Patent No. 5405942
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#### ALIGNMENTS

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RESULT 1
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; Sequence 3, Application US/09142583A
 Patent No. 6221842
    GENERAL INFORMATION:
         APPLICANT: GOLDSPINK, GEOFFREY
         TITLE OF INVENTION: METHOD OF TREATING MUSCULAR DISORDERS
;
         NUMBER OF SEQUENCES: 11
;
         CORRESPONDENCE ADDRESS:
              ADDRESSEE: NIXON & VANDERHYE P.C.
              STREET: 1100 NORTH GLEBE ROAD
              CITY: ARLINGTON
              STATE: VA
              COUNTRY: USA
              ZIP: 22201
         COMPUTER READABLE FORM:
              MEDIUM TYPE: Floppy disk
              COMPUTER: IBM PC compatible
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OPERATING SYSTEM: PC-DOS/MS-DOS
          SOFTWARE: PatentIn Release #1.0, Version #1.25
      CURRENT APPLICATION DATA:
          APPLICATION NUMBER: US/09/142,583A
          FILING DATE: 29-Oct-1998
          CLASSIFICATION: <Unknown>
       PRIOR APPLICATION DATA:
          APPLICATION NUMBER: WO PCT/GB97/00658
          FILING DATE: 11-MAR-1997
          APPLICATION NUMBER: GB 9605124.8
          FILING DATE: 11-MAR-1996
      ATTORNEY/AGENT INFORMATION:
          NAME: SADOFF, B. J.
          REGISTRATION NUMBER: 36663
          REFERENCE/DOCKET NUMBER: 117-263
      TELECOMMUNICATION INFORMATION:
          TELEPHONE: 7038164000
          TELEFAX: 7038164100
   INFORMATION FOR SEQ ID NO: 3:
      SEQUENCE CHARACTERISTICS:
          LENGTH: 553 base pairs
          TYPE: nucleic acid
          STRANDEDNESS: both
          TOPOLOGY: linear
      MOLECULE TYPE: cDNA
      FEATURE:
          NAME/KEY: CDS
          LOCATION: 1..363
      SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-09-142-583A-3
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                          Pred. No. 1.6e-154;
 Best Local Similarity
                   100.0%;
 Matches 523; Conservative
                         0; Mismatches
                                       0; Indels
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US-09-142-583A-5
; Sequence 5, Application US/09142583A
; Patent No. 6221842
   GENERAL INFORMATION:
        APPLICANT: GOLDSPINK, GEOFFREY
        TITLE OF INVENTION: METHOD OF TREATING MUSCULAR DISORDERS
        NUMBER OF SEQUENCES: 11
        CORRESPONDENCE ADDRESS:
            ADDRESSEE: NIXON & VANDERHYE P.C.
            STREET: 1100 NORTH GLEBE ROAD
            CITY: ARLINGTON
            STATE: VA
            COUNTRY: USA
            ZIP: 22201
        COMPUTER READABLE FORM:
            MEDIUM TYPE: Floppy disk
            COMPUTER: IBM PC compatible
            OPERATING SYSTEM: PC-DOS/MS-DOS
            SOFTWARE: PatentIn Release #1.0, Version #1.25
        CURRENT APPLICATION DATA:
            APPLICATION NUMBER: US/09/142,583A
            FILING DATE: 29-Oct-1998
            CLASSIFICATION: <Unknown>
        PRIOR APPLICATION DATA:
            APPLICATION NUMBER: WO PCT/GB97/00658
            FILING DATE: 11-MAR-1997
            APPLICATION NUMBER: GB 9605124.8
            FILING DATE: 11-MAR-1996
        ATTORNEY/AGENT INFORMATION:
            NAME: SADOFF, B. J.
            REGISTRATION NUMBER: 36663
            REFERENCE/DOCKET NUMBER: 117-263
        TELECOMMUNICATION INFORMATION:
            TELEPHONE: 7038164000
            TELEFAX: 7038164100
   INFORMATION FOR SEQ ID NO: 5:
        SEQUENCE CHARACTERISTICS:
            LENGTH: 553 base pairs
            TYPE: nucleic acid
            STRANDEDNESS: both
            TOPOLOGY: linear
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MOLECULE TYPE: cDNA
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          NAME/KEY: CDS
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                        Score 523; DB 3; Length 553;
 Best Local Similarity
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; Sequence 10, Application US/09142583A
 Patent No. 6221842
  GENERAL INFORMATION:
      APPLICANT: GOLDSPINK, GEOFFREY
      TITLE OF INVENTION: METHOD OF TREATING MUSCULAR DISORDERS
      NUMBER OF SEQUENCES: 11
      CORRESPONDENCE ADDRESS:
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ADDRESSEE: NIXON & VANDERHYE P.C.
            STREET: 1100 NORTH GLEBE ROAD
            CITY: ARLINGTON
            STATE: VA
            COUNTRY: USA
            ZIP: 22201
       COMPUTER READABLE FORM:
            MEDIUM TYPE: Floppy disk
            COMPUTER: IBM PC compatible
            OPERATING SYSTEM: PC-DOS/MS-DOS
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            FILING DATE: 29-Oct-1998
            CLASSIFICATION: <Unknown>
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            APPLICATION NUMBER: WO PCT/GB97/00658
            FILING DATE: 11-MAR-1997
            APPLICATION NUMBER: GB 9605124.8
            FILING DATE: 11-MAR-1996
       ATTORNEY/AGENT INFORMATION:
            NAME: SADOFF, B. J.
            REGISTRATION NUMBER: 36663
            REFERENCE/DOCKET NUMBER: 117-263
       TELECOMMUNICATION INFORMATION:
            TELEPHONE: 7038164000
            TELEFAX: 7038164100
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            TOPOLOGY: linear
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    APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER,
; JAMES P.
   TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
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   NUMBER OF SEQUENCES: 16
    CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/07/65,673
     FILING DATE: 16-JUN-1987
    PRIOR APPLICATION DATA:
     APPLICATION NUMBER: 630,557
     FILING DATE: 19-JUL-1984
;SEQ ID NO:2:
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RESULT 5
US-08-472-809B-8
; Sequence 8, Application US/08472809B
 Patent No. 5925564
  GENERAL INFORMATION:
   APPLICANT: Schwartz, Robert J.
   APPLICANT: DeMayo, Franco J.
   APPLICANT: O'Malley, Bert W.
   TITLE OF INVENTION: Expression Vector Systems and
   TITLE OF INVENTION: Method of Use
   NUMBER OF SEQUENCES: 8
   CORRESPONDENCE ADDRESS:
   ADDRESSEE: Lyon & Lyon
     STREET: 633 West Fifth Street
     STREET: Suite 4700
     CITY: Los Angeles
     STATE: California
     COUNTRY: U.S.A.
     ZIP: 90071-2066
   COMPUTER READABLE FORM:
     MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
     MEDIUM TYPE: storage
     COMPUTER: IBM Compatible
     OPERATING SYSTEM: IBM P.C. DOS 5.0
     SOFTWARE: Word Perfect 5.1
   CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/08/472,809B
     FILING DATE: June 7, 1995
     CLASSIFICATION: 435
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PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/209,846

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FILING DATE: March 9, 1994
     APPLICATION NUMBER: 07/789,919
     FILING DATE: No. 5925564ember 6, 1991
    ATTORNEY/AGENT INFORMATION:
     NAME: Warburg, Richard J.
     REGISTRATION NUMBER: 32,327
     REFERENCE/DOCKET NUMBER: 214/212
    TELECOMMUNICATION INFORMATION:
     TELEPHONE: (213) 489-1600
     TELEFAX: (213) 955-0440
     TELEX: 67-3510
  INFORMATION FOR SEQ ID NO: 8:
    SEQUENCE CHARACTERISTICS:
     LENGTH: 5707 bases
     TYPE: nucleic acid
     STRANDEDNESS: double
     TOPOLOGY: linear
   MOLECULE TYPE: cDNA
US-08-472-809B-8
 Query Match
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 Best Local Similarity 82.2%; Pred. No. 5.3e-76;
 Matches 351; Conservative 0; Mismatches 24; Indels
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RESULT 6
US-08-472-809B-7
; Sequence 7, Application US/08472809B
; Patent No. 5925564
   GENERAL INFORMATION:
     APPLICANT: Schwartz, Robert J.
    APPLICANT: DeMayo, Franco J.
    APPLICANT: O'Malley, Bert W.
    TITLE OF INVENTION: Expression Vector Systems and
    TITLE OF INVENTION: Method of Use
    NUMBER OF SEQUENCES: 8
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Lyon & Lyon
      STREET: 633 West Fifth Street
      STREET: Suite 4700
      CITY: Los Angeles
      STATE: California
      COUNTRY: U.S.A.
      ZIP: 90071-2066
    COMPUTER READABLE FORM:
      MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
      MEDIUM TYPE: storage
      COMPUTER: IBM Compatible
      OPERATING SYSTEM: IBM P.C. DOS 5.0
      SOFTWARE: Word Perfect 5.1
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/472,809B
      FILING DATE: June 7, 1995
      CLASSIFICATION: 435
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: 08/209,846
      FILING DATE: March 9, 1994
      APPLICATION NUMBER: 07/789,919
      FILING DATE: No. 5925564ember 6, 1991
    ATTORNEY/AGENT INFORMATION:
      NAME: Warburg, Richard J.
      REGISTRATION NUMBER: 32,327
      REFERENCE/DOCKET NUMBER:
                               214/212
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (213) 489-1600
      TELEFAX: (213) 955-0440
;
      TELEX: 67-3510
  INFORMATION FOR SEQ ID NO: 7:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6345 bases
      TYPE: nucleic acid
      STRANDEDNESS: double
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
US-08-472-809B-7
 Query Match
                        52.5%; Score 274.6; DB 2; Length 6345;
 Best Local Similarity 82.2%; Pred. No. 5.6e-76;
 Matches 351; Conservative 0; Mismatches
                                              24; Indels
                                                            52; Gaps
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; Patent No. 5405942
   APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER,
   TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
;I AND II
   NUMBER OF SEQUENCES: 16
   CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/07/65,673
     FILING DATE: 16-JUN-1987
   PRIOR APPLICATION DATA:
     APPLICATION NUMBER: 630,557
     FILING DATE: 19-JUL-1984
;SEQ ID NO:13:
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RESULT 8
5405942-9
; Patent No. 5405942
   APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER.
   TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
;I AND II
   NUMBER OF SEQUENCES: 16
   CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/07/65,673
     FILING DATE: 16-JUN-1987
   PRIOR APPLICATION DATA:
     APPLICATION NUMBER: 630,557
     FILING DATE: 19-JUL-1984
;SEQ ID NO:9:
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    APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER,
; JAMES P.
    TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
; I AND II
    NUMBER OF SEQUENCES: 16
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/07/65,673
      FILING DATE: 16-JUN-1987
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: 630,557
      FILING DATE: 19-JUL-1984
;SEO ID NO:7:
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RESULT 10
5405942-11
; Patent No. 5405942
    APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER,
; JAMES P.
    TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
;I AND II
    NUMBER OF SEQUENCES: 16
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/07/65,673
      FILING DATE: 16-JUN-1987
    PRIOR APPLICATION DATA:
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      FILING DATE: 19-JUL-1984
; SEQ ID NO:11:
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RESULT 11
US-09-255-829-13
; Sequence 13, Application US/09255829
; Patent No. 6461617
  GENERAL INFORMATION:
    APPLICANT: Shone, Clifford Charles
    APPLICANT: Quinn, Conrad Padraig
APPLICANT: Foster, Keith Alan
    TITLE OF INVENTION: Recombinant Toxin Fragments
    NUMBER OF SEQUENCES: 29
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: STERNE, KESSLER, GOLDSTEIN, & FOX P.L.L.C.
      STREET: 1100 NEW YORK AVENUE, NW, SUITE 600
      CITY: WASHINGTON
      STATE: DC
      COUNTRY: USA
      ZIP: 20005-3934
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)
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      APPLICATION NUMBER: PCT/GB97/02273
      FILING DATE: 22-AUG-1997
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 08/782,893
      FILING DATE: 27-DEC-1996
    ATTORNEY/AGENT INFORMATION:
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NAME: ESMOND, ROBERT W.

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REGISTRATION NUMBER: 32,893
      REFERENCE/DOCKET NUMBER: 1581.0130002
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 202-371-2600
      TELEFAX: 202-371-2540
  INFORMATION FOR SEQ ID NO: 13:
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      STRANDEDNESS: double
      TOPOLOGY: linear
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    APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER,
; JAMES P.
    TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
; I AND II
    NUMBER OF SEQUENCES: 16
    CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/07/65,673
     FILING DATE: 16-JUN-1987
    PRIOR APPLICATION DATA:
     APPLICATION NUMBER: 630,557
     FILING DATE: 19-JUL-1984
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; Sequence 1, Application US/08308196A
; Patent No. 5612198
  GENERAL INFORMATION:
    APPLICANT: Brierley, Russell A.
    APPLICANT: Davis, Geneva R.
    APPLICANT: Holtz, Gregory C.
    APPLICANT: Gleeson, Martin A.
    APPLICANT: Howard, Bradley D.
    TITLE OF INVENTION: Production of Insulin-Like Growth
TITLE OF INVENTION: Factor-1 in Methylotrophic Yeast Cells
    NUMBER OF SEQUENCES: 17
    CORRESPONDENCE ADDRESS:
     ADDRESSEE: Brown, Martin, Haller & McClain
      STREET: 1660 Union Street
      CITY: San Diego
      STATE: California
      COUNTRY: USA
      ZIP: 92101-2926
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/308,196A
      FILING DATE: 09-SEPT-1994
    PRIOR APPLICATION DATA:
     APPLICATION NUMBER: US 07/983,523
      FILING DATE: 03-MAR-1993
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 07/578,728
      FILING DATE: 04-SEP-1990
    ATTORNEY/AGENT INFORMATION:
     NAME: Seidman, Stephanie L.
      REGISTRATION NUMBER: 33,779
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REFERENCE/DOCKET NUMBER:
                             51875
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (619)238-0999
      TELEFAX: (619)238-0062
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 240 base pairs
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    APPLICANT: Brierley, Russell A.
    APPLICANT: Davis, Geneva R.
    APPLICANT: Holtz, Gregory C.
    APPLICANT: Gleeson, Martin A.
    APPLICANT: Bradley, D. H.
    TITLE OF INVENTION: Production of Insulin-Like Growth
    TITLE OF INVENTION: Factor-1 in Methylotrophic Yeast Cells
    NUMBER OF SEQUENCES: 12
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Fitch, Even, Tabin & Flannery
      STREET: 135 South LaSalle Street, Suite 900
      CITY: Chicago
      STATE: Illinois
     COUNTRY: USA
      ZIP: 60603
    COMPUTER READABLE FORM:
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      FILING DATE: 19910409
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      FILING DATE: 04-SEP-1990
    ATTORNEY/AGENT INFORMATION:
      NAME: Seidman, Stephanie L.
      REGISTRATION NUMBER: 33,779
      REFERENCE/DOCKET NUMBER:
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (619)552-1311
      TELEFAX: (619)552-0095
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; Patent No. 6107057
  GENERAL INFORMATION:
    APPLICANT: Crawford, Kenneth
    APPLICANT: Zaror, Isabel
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APPLICANT: Innis, Michael
    TITLE OF INVENTION: Pichia Secretory Leader for Protein
    TITLE OF INVENTION: Expression
    NUMBER OF SEQUENCES: 40
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Chiron Corporation
      STREET: 4560 Horton Street
      CITY: Emeryville
      STATE: California
      COUNTRY: United States
      ZIP: 94608
    COMPUTER READABLE FORM:
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    ATTORNEY/AGENT INFORMATION:
      NAME: Chung, Ling-Fong
      REGISTRATION NUMBER: 36,482
      REFERENCE/DOCKET NUMBER: 1165.100
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (510) 601-2704
      TELEFAX: (510) 655-3542
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US-09-852-261-5 Title:

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Post-processing: Minimum Match 0%

Maximum Match 100%

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score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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> No. Score Match Length DB ID

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	8	334.4	63.9	7260	13	US-09-873-319-707	Sequence 707, App
	9	334.4	63.9	7260	13	US-09-960-706-1066	Sequence 1066, Ap
	10	334.4	63.9	7260	15	US-10-136-639-4	Sequence 4, Appli
	11	332.8	63.6	725	15	US-10-207-655-54	Sequence 54, Appl
	12	273.6	52.3	612	13	US-10-251-661-7	Sequence 7, Appli
	13	262	50.1	487	9	US-09-852-261-11	Sequence 11, Appl
	14	237.6	45.4	318	9	US-09-852-261-9	Sequence 9, Appli
	15	228	43.6	462	15	US-10-238-114-1	Sequence 1, Appli
	16	209	40.0	286	15	US-10-161-088-3	Sequence 3, Appli
	17	193	36.9	210	13	US-09-807-742-18	Sequence 18, Appl
	18	191.4	36.6	2862	13	US-10-241-596-13	Sequence 13, Appl
	19	187	35.8	4532	10	US-09-930-377B-1	Sequence 1, Appli
	20	186.6	35.7	210	10	US-09-930-377B-2	Sequence 2, Appli
	21	185.2	35.4	390	15	US-10-179-046-13	Sequence 13, Appl
	22	163.8	31.3	516	13	US-10-029-386-5832	Sequence 5832, Ap
	23	162.8	31.1	182	13	US-10-029-386-18231	Sequence 18231, A
	24	141.6	27.1	213	15	US-10-076-816-9	Sequence 9, Appli
	25	141.6	27.1	213	15	US-10-077-381-9	Sequence 9, Appli
	26	127	24.3	621	9	US-09-921-398-40	Sequence 40, Appl
	27	·127	24.3	621	15	US-10-280-826-40	Sequence 40, Appl
	28	113.4	21.7	480	9	US-09-921-398-38	Sequence 38, Appl
	29	113.4	21.7	480	15	US-10-280-826-38	Sequence 38, Appl
	30	101.8	19.5	210	13	US-09-807-742-19	Sequence 19, Appl
	31	77.2	14.8	854	10	US-09-954-531-989	Sequence 989, App
С	32	75.4	14.4	447	9	US-09-922-217-917	Sequence 917, App
С	33	75.4	14.4	447	10	US-09-833-263-917	Sequence 917, App
С	34	75.4	14.4	447	14	US-10-025-380-917	Sequence 917, App
С	35	75.2	14.4	437	15	US-10-066-543-663	Sequence 663, App
С	36	75.2	14.4	493	15	US-10-066-543-997	Sequence 997, App
С	37	75.2	14.4	518	15	US-10-066-543-1040	Sequence 1040, Ap
С	38	75.2	14.4	536	15	US-10-066-543-428	Sequence 428, App
	39	75.2	14.4	543	15	US-10-136-841-1	Sequence 1, Appli
С	40	75.2	14.4	549	15	US-10-066-543-478	Sequence 478, App
С	41	75.2	14.4	574	9	US-09-922-217-918	Sequence 918, App
С	42	75.2	14.4	574	10	US-09-833-263-918	Sequence 918, App
С	43	75.2	14.4	574	14	US-10-025-380-918	Sequence 918, App
С	44	75.2	14.4	577	15	US-10-066-543-1137	Sequence 1137, Ap
С	45	75.2	14.4	579	15	US-10-066-543-1094	Sequence 1094, Ap

## ALIGNMENTS

# RESULT 1

US-09-852-261-5

- ; Sequence 5, Application US/09852261
  ; Patent No. US20020083477A1
- ; GENERAL INFORMATION:
- ; APPLICANT: GOLDSPINK, GEOFFREY

```
APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 5
  LENGTH: 523
  TYPE: DNA
  ORGANISM: Oryctolagus cuniculus
US-09-852-261-5
 Query Match
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 Best Local Similarity
                  100.0%; Pred. No. 1.9e-161;
                       0: Mismatches
 Matches 523; Conservative
                                    0: Indels
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                                                 Gaps
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          1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
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          61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
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          Db
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       Qу
          Db
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          Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
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          301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
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Qу
          Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
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          Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
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Qу
          481 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Db
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US-09-852-261-1
; Sequence 1, Application US/09852261
; Patent No. US20020083477A1
 GENERAL INFORMATION:
  APPLICANT: GOLDSPINK, GEOFFREY
  APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
   LENGTH: 517
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-852-261-1
 Query Match
                    89.4%; Score 467.4; DB 9; Length 517;
 Best Local Similarity 96.2%; Pred. No. 3.8e-143;
 Matches 501; Conservative
                         0; Mismatches 16;
                                          Indels
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           61 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 120
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       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
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           241 ATGCCCAAGACCCAGAAGTATCAGCCCCCATCTACCAACAAGAACACGAAGTCTCA---G 297
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           298 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 357
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QУ
           Db
       358 CAGGATGTA-GAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 416
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           Db
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Db

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RESULT 3
US-09-852-261-13
; Sequence 13, Application US/09852261
; Patent No. US20020083477A1
; GENERAL INFORMATION:
  APPLICANT: GOLDSPINK, GEOFFREY
  APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 13
  LENGTH: 471
  TYPE: DNA
  ORGANISM: Oryctolagus cuniculus
US-09-852-261-13
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 Best Local Similarity 90.1%; Pred. No. 6.1e-124;
 Matches 471; Conservative
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          121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
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       Qy
          Db
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Db
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           429 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 471
Db
RESULT 4
US-09-852-261-3
; Sequence 3, Application US/09852261
; Patent No. US20020083477A1
; GENERAL INFORMATION:
  APPLICANT: GOLDSPINK, GEOFFREY
  APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 3
   LENGTH: 539
   TYPE: DNA
   ORGANISM: Rattus sp.
US-09-852-261-3
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                                          Length 539;
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 Best Local Similarity
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 Matches 436; Conservative
                                           Indels
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           481 TCATTTCAGAGATGGGCATTTCCCTCAATGAAATACACAAGTAAACATTC 530
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RESULT 5
US-10-161-088-1
; Sequence 1, Application US/10161088
; Publication No. US20030077761A1
; GENERAL INFORMATION:
  APPLICANT: Parrow, Vendela
  APPLICANT: Rosengren, Linda
  TITLE OF INVENTION: NEW METHODS
  FILE REFERENCE: 13425-111001
  CURRENT APPLICATION NUMBER: US/10/161,088
  CURRENT FILING DATE: 2002-05-31
  PRIOR APPLICATION NUMBER: SE 0101934-8
  PRIOR FILING DATE: 2001-06-01
  NUMBER OF SEO ID NOS: 3
  SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 1
   LENGTH: 651
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (73)...(471)
US-10-161-088-1
                    66.8%; Score 349.4; DB 15; Length 651;
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 Matches 425; Conservative
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Qу
           1111
                                   499 CAGAATGTAGGAGGAGCCTCCCACGGAGCAGAAAATGCCACATCACCGCAGGATCCTTTG 558
Db
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                     1
                         \mathbf{I}
        559 CTGCTTGAGCAACCTGCAAAACATCGAAACACCTACCAAATAACAATAATAAGTCCAATA 618
Db
        475 ACATTTCAAAGAT-GGCATTTCCCCCAATGAAA 506
Qy
           619 ACATTACAAAGATGGGCATTTCCCCCAATGAAA 651
Db
RESULT 6
US-09-919-497-24
; Sequence 24, Application US/09919497
; Patent No. US20020106662A1
; GENERAL INFORMATION:
  APPLICANT: Mutter, George L.
  TITLE OF INVENTION: PROGNOSTIC CLASSIFICATION OF ENDOMETRIAL CANCER
  FILE REFERENCE: B0801/7225
  CURRENT APPLICATION NUMBER: US/09/919,497
  CURRENT FILING DATE: 2001-07-31
  PRIOR APPLICATION NUMBER: US 60/221,735
  PRIOR FILING DATE: 2000-07-31
  NUMBER OF SEQ ID NOS: 100
  SOFTWARE: PatentIn version 3.0
 SEQ ID NO 24
   LENGTH: 7260
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-919-497-24
                           Score 334.4; DB 10; Length 7260;
                    63.9%;
 Query Match
 Best Local Similarity
                    84.6%; Pred. No. 9e-99;
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 Matches 445; Conservative
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Qy
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           371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
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        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
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           431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
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           551 ATGCCCAAGACCCAG----- 565
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QУ
                Db
        566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
        361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qy
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
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           679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
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        479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           739 TTAAAAGATGGGCGTTTCCCCCCAATGAAATACACAAGTAAACATTC 784
Db
RESULT 7
US-09-880-107-3739
; Sequence 3739, Application US/09880107
; Patent No. US20020142981A1
; GENERAL INFORMATION:
  APPLICANT: Horne, Darci T.
  APPLICANT: Vockley, Joseph G.
; APPLICANT: Scherf, Uwe
 APPLICANT: Gene Logic, Inc.
  TITLE OF INVENTION: Gene Expression Profiles in Liver Cancer
  FILE REFERENCE: 44921-5028-WO
  CURRENT APPLICATION NUMBER: US/09/880,107
  CURRENT FILING DATE: 2001-06-14
  PRIOR APPLICATION NUMBER: US 60/211,379
  PRIOR FILING DATE: 2000-06-14
  PRIOR APPLICATION NUMBER: US 60/237,054
  PRIOR FILING DATE: 2000-10-02
  NUMBER OF SEQ ID NOS: 3950
  SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3739
  LENGTH: 7260
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   OTHER INFORMATION: Genbank Accession No. US20020142981A1 X57025
US-09-880-107-3739
 Query Match
                    63.9%; Score 334.4; DB 10; Length 7260;
 Best Local Similarity 84.6%; Pred. No. 9e-99;
 Matches 445; Conservative 0; Mismatches 26; Indels
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Db	431	${\tt ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT}$	490
Qу	181	TGTGCACCCTCAAGCCGGCAAAGGCAGCCCGCTCCGTCCG	240
Db	491	TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC	550
Qу	241	ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG	300
Db	551	ATGCCCAAGACCCAG	565
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Db	566	AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA	618
QУ	361	CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG	420
Db	619	CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG	678
QУ	421	CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT	478
Db	679		738
QУ	479	TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523	
Db	739	TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 784	

## RESULT 8

US-09-873-319-707

- ; Sequence 707, Application US/09873319A
- ; Publication No. US20030134324A1
- ; GENERAL INFORMATION:
- ; APPLICANT: Munger, William E.
- ; APPLICANT: Kulkarni, Prakash
- ; APPLICANT: Getzenberg, Robert H.
- ; APPLICANT: Waga, Iwao
- ; APPLICANT: Yamamoto, Jun
- ; TITLE OF INVENTION: Identifying Drugs for and Diagnosis of Benign Prostatic
- ; TITLE OF INVENTION: Hyperplasia Using Gene Expression Profiles
- ; FILE REFERENCE: 44921-5029-US
- ; CURRENT APPLICATION NUMBER: US/09/873,319A
- ; CURRENT FILING DATE: 2001-06-05
- ; EARLIER APPLICATION NUMBER: US 60/223,323
- ; EARLIER FILING DATE: 2000-08-07
- ; NUMBER OF SEQ ID NOS: 755
- ; SOFTWARE: PatentIn Ver. 2.1
- ; SEQ ID NO 707

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LENGTH: 7260
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   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   OTHER INFORMATION: Genbank Accession No. US20030134324A1 X57025
US-09-873-319-707
                   63.9%; Score 334.4; DB 13; Length 7260;
 Query Match
 Best Local Similarity 84.6%; Pred. No. 9e-99;
                         0; Mismatches 26; Indels
                                                55; Gaps
 Matches 445; Conservative
                                                          4;
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGAGAC 60
Qу
          311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
          371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
          431 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
       Qу
          491 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 550
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qy
          Db
       551 ATGCCCAAGACCCAG----
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qy
                566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qy
          619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
       421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qy
          679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
Db
       479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qy
          739 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 784
Db
RESULT 9
US-09-960-706-1066
; Sequence 1066, Application US/09960706
; Publication No. US20030134280A1
; GENERAL INFORMATION:
 APPLICANT: Munger, William E.
  TITLE OF INVENTION: Identifying Drugs for and Diagnosis of Benign Prostatic
Hyperplasia Using
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; TITLE OF INVENTION: Gene Expression Profiles

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FILE REFERENCE: 44921-5029-01US
  CURRENT APPLICATION NUMBER: US/09/960,706
  CURRENT FILING DATE: 2001-09-24
  PRIOR APPLICATION NUMBER: 60/223,323
  PRIOR FILING DATE: 2000-08-07
  PRIOR APPLICATION NUMBER: 09/873,319
  PRIOR FILING DATE: 2001-06-05
  NUMBER OF SEQ ID NOS: 1124
  SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 1066
   LENGTH: 7260
   TYPE: DNA
   ORGANISM: Homo sapiens
   OTHER INFORMATION: Genbank Accession No. US20030134280A1 X57025
US-09-960-706-1066
 Query Match
                   63.9%; Score 334.4; DB 13; Length 7260;
 Best Local Similarity 84.6%; Pred. No. 9e-99;
 Matches 445; Conservative
                         0; Mismatches
                                      26;
                                          Indels
                                                    Gaps
                                                          4;
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qy
          311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
0v
          371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qy
          431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
       Qy
          491 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 550
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qν
           1111111111 111
       551 ATGCCCAAGACCCAG----- 565
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
                566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
       421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
           679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
Db
       479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           Db
       739 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 784
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RESULT 10
US-10-136-639-4
; Sequence 4, Application US/10136639
; Publication No. US20030072761A1
; GENERAL INFORMATION:
  APPLICANT: LeBowitz, Jonathan
  TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR TARGETING PROTEINS ACROSS
THE BLOOD BRAIN
  TITLE OF INVENTION: BARRIER
  FILE REFERENCE: SYM-008
  CURRENT APPLICATION NUMBER: US/10/136,639
  CURRENT FILING DATE: 2002-09-06
  PRIOR APPLICATION NUMBER: US 60/329,650
  PRIOR FILING DATE: 2001-10-16
  NUMBER OF SEQ ID NOS: 4
  SOFTWARE: PatentIn version 3.0
SEO ID NO 4
   LENGTH: 7260
   TYPE: DNA
   ORGANISM: Homo sapiens
US-10-136-639-4
                   63.9%; Score 334.4; DB 15; Length 7260;
 Query Match
 Best Local Similarity 84.6%; Pred. No. 9e-99;
                                                          4;
 Matches 445; Conservative
                         0; Mismatches 26; Indels
                                                55;
                                                    Gaps
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Qу
          311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 370
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
          371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
          431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
       Qу
          491 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 550
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
          551 ATGCCCAAGACCCAG----- 565
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
               566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
       421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
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Db
       679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
       479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           739 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 784
Db
RESULT 11
US-10-207-655-54
; Sequence 54, Application US/10207655
; Publication No. US20030118592A1
; GENERAL INFORMATION:
  APPLICANT: Ledbetter, Jeffrey A.
  APPLICANT: Hayden-Ledbetter, Martha S.
  TITLE OF INVENTION: BINDING DOMAIN-IMMUNOGLOBULIN FUSION PROTEINS
  FILE REFERENCE: 390069.401C1
  CURRENT APPLICATION NUMBER: US/10/207,655
  CURRENT FILING DATE: 2002-07-25
  NUMBER OF SEO ID NOS: 426
  SOFTWARE: PatentIn version 3.0
; SEQ ID NO 54
   LENGTH: 725
   TYPE: DNA
   ORGANISM: Homo sapiens
US-10-207-655-54
                    63.6%; Score 332.8; DB 15; Length 725;
 Query Match
 Best Local Similarity 84.4%; Pred. No. 9.2e-99;
                         0: Mismatches '27: Indels
 Matches 444; Conservative
                                                            4:
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qy
           156 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 215
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
           216 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 275
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qy
           276 ACAGGTATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 335
Db
       Qy
           336 TGCGCACCCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 395
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qy
           11[1]11111 111
       396 ATGCCCAAGACCCAG----- 410
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qy
                411 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 463
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qy
           464 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 523
Db
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421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
           524 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 583
Db
        479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           Db
        584 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 629
RESULT 12
US-10-251-661-7
; Sequence 7, Application US/10251661
; Publication No. US20030166555A1
; GENERAL INFORMATION:
  APPLICANT: Alberini, Cristina M.
  APPLICANT: Bear, Mark F.
  TITLE OF INVENTION: Methods and Compositions for Regulating
  TITLE OF INVENTION: Memory Consolidation
  FILE REFERENCE: 3499.1001-003
  CURRENT APPLICATION NUMBER: US/10/251,661
  CURRENT FILING DATE: 2002-09-20
  PRIOR APPLICATION NUMBER: 60/193,614
  PRIOR FILING DATE: 2000-03-31
  PRIOR APPLICATION NUMBER: PCT/US01/10661
  PRIOR FILING DATE: 2001-04-02
  NUMBER OF SEQ ID NOS: 12
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEO ID NO 7
   LENGTH: 612
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (151)...(564)
US-10-251-661-7
 Query Match
                    52.3%; Score 273.6; DB 13; Length 612;
 Best Local Similarity 83.0%; Pred. No. 2.6e-79;
 Matches 347; Conservative
                         0; Mismatches 19; Indels
                                                             1:
                                                   52; Gaps
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Qy
           Db
       247 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 306
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qy
           Db
        307 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 366
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qy
           367 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 426
Db
        Qy
           427 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 486
Db
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241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
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Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
                502 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 554
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTT 418
QУ
          555 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTT 612
Db
RESULT 13
US-09-852-261-11
; Sequence 11, Application US/09852261
; Patent No. US20020083477A1
; GENERAL INFORMATION:
 APPLICANT: GOLDSPINK, GEOFFREY
 APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
; SEO ID NO 11
   LENGTH: 487
   TYPE: DNA
   ORGANISM: Rattus sp.
US-09-852-261-11
                   50.1%; Score 262; DB 9; Length 487;
 Query Match
 Best Local Similarity 74.7%; Pred. No. 1.5e-75;
 Matches 396; Conservative 0; Mismatches 75; Indels
                                                 59; Gaps
                                                           3:
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
          1 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGACGCTCTTCAGTTCGTGTGTGGACCA 60
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
          61 AGGGGCTTTTACTTCAACAAGCCCACAGTCTATGGCTCCAGCATTCGGAGGGCACCACAG 120
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
          121 ACGGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
       Qу
          181 TGTGTCCGCTGCAAGCCTACAAAGTCAGCTCGTTCCATCCGGGCCCAGCGCCACACTGAC 240
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qy
          241 ATGCCCAAGACTCAG----- 255
Db
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301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
                256 -----AAGGAAGTACACTTGAAGAACACAAGTAGAGGAAGTGCAGGAAACAAGACCTA 308
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
                                   111 1111111 11 111 1 11111
        309 CAGAATGTAGGAGGAGCCTCCCGAGGAACAGAAAATGCCACGTCACCGCAAGATCCTTTG 368
Db
        421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA-----AAAAATAAGTTTGATC 474
Qу
                     1
                         11111 1111 111
                                              11 111 1111 11
           369 CTGCTTGAGCAACCTGCAAAACATCGGAACACCTGCCAAATATCAATAATGAGTTCAATA 428
Db
        475 ACATTTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
            429 TCATTTCAGAGATGGGCATTTCCCTCAATGAAATACACAAGTAAACATTC 478
Db
RESULT 14
US-09-852-261-9
; Sequence 9, Application US/09852261
; Patent No. US20020083477A1
; GENERAL INFORMATION:
  APPLICANT: GOLDSPINK, GEOFFREY
  APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 9
   LENGTH: 318
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-852-261-9
                     45.4%; Score 237.6; DB 9; Length 318;
 Query Match
                          Pred. No. 1.3e-67;
                    94.6%;
 Best Local Similarity
                                                             0;
                          0; Mismatches 14;
                                            Indels
 Matches 246; Conservative
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGAGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qy
           1 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Db
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
QУ
           61 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 120
Db
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           121 ACAGGCATCGTGGATGACTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 180
Db
        Qy
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181 TGCGCACCCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 240
Db
        241 ATGCCCAAGACTCAGAAGTA 260
Qy
           241 ATGCCCAAGACCCAGAAGGA 260
Db
RESULT 15
US-10-238-114-1
; Sequence 1, Application US/10238114
; Publication No. US20030100073A1
; GENERAL INFORMATION:
; APPLICANT: Merial
; APPLICANT: ANDREONI , Christine Michele
  TITLE OF INVENTION: IGF-1 AS FELINE VACCINE ADJUVANT, IN PARTICULAR AGAINST
FELINE RETROVIRUS
  FILE REFERENCE: 454313-3165.1
  CURRENT APPLICATION NUMBER: US/10/238,114
  CURRENT FILING DATE: 2002-09-10
  PRIOR APPLICATION NUMBER: FR 01 11736
  PRIOR FILING DATE: 2001-09-11
  PRIOR APPLICATION NUMBER: US 60/318,666
  PRIOR FILING DATE: 2001-09-12
  NUMBER OF SEQ ID NOS: 20
  SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
   LENGTH: 462
   TYPE: DNA
   ORGANISM: Felis catus
US-10-238-114-1
                     43.6%; Score 228; DB 15; Length 462;
 Query Match
 Best Local Similarity 92.3%; Pred. No. 2.3e-64;
                                                      0; Gaps
                                         20; Indels
                                                                0;
 Matches 240; Conservative 0; Mismatches
          1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGAGACCTCTTCAGTTCGTGTGTGGAGAC 60
Qy
            145 GGACCAGAGACGCTCTGTGGGGCTGAGTTGGTGGACGCTCTTCAGTTCGTGTGGAGAC 204
Db
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qy
           205 AGGGGTTTTTATTTCAACAAGCCCACGGGGTATGGCTCCAGCAGTCGGAGGGCACCTCAG 264
Db
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qy
            265 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGCGGCTAGAGATGTAC 324
Db
        Qy
            325 TGTGCACCCCTCAAGCCTGCCAAGTCTGCCCGCTCAGTCCGTGCTCAGCGCCACACTGAC 384
Db
        241 ATGCCCAAGACTCAGAAGTA 260
Qy
            11111111 11111
        385 ATGCCCAAGGCTCAGAAGGA 404
Db
Search completed: December 13, 2003, 11:56:48
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Job time : 235.512 secs